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| (54) Title: <b>A NOVEL POLYMORPHIC CRYSTALLINE FORM OF FLUTICASONE PROPIONATE, A METHOD FOR ITS PRODUCTION AND PHARMACEUTICAL COMPOSITIONS THEREOF</b>  |  |  |   |
| (57) Abstract<br><p>The invention relates to a new polymorphic crystalline form of S-fluoromethyl 6<math>\alpha</math>,9<math>\alpha</math>-difluoro-11<math>\beta</math>-hydroxy-16<math>\alpha</math>-methyl-3-oxo-17<math>\alpha</math>-propionyloxyandrosta-1,4-diene-17<math>\beta</math>-carbothiate (fluticasone propionate). The new polymorphic crystalline form is easily handled and easily fluidised and its particle size and shape can be controlled. The invention also relates to the use of this new material in therapy, particularly in the treatment of respiratory disorders, e.g. asthma.</p> |  |  |   |

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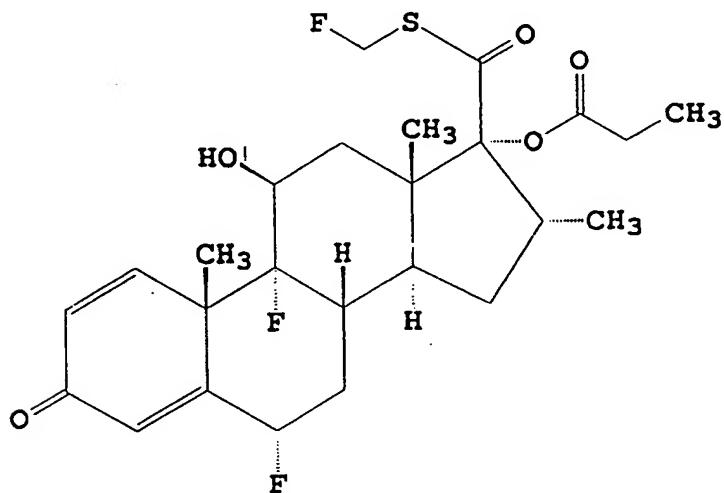
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A METHOD FOR ITS PRODUCTION AND PHARMACEUTICAL COMPOSITIONS THEREOF  
A NOVEL POLYMORPHIC CRYSTALLINE FORM OF FLUTICASONE PROPIONATE,

5 The present invention relates to particulate products  
which may be prepared by using supercritical fluids.  
More particularly, the invention relates to novel  
crystalline forms of fluticasone propionate, which is  
10 S-fluoromethyl 6 $\alpha$ ,9 $\alpha$ -difluoro-11 $\beta$ -hydroxy-16 $\alpha$ -methyl-3-  
oxo-17 $\alpha$ -propionyloxyandrosta-1,4-diene-17 $\beta$ -carbothiate.

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Fluticasone propionate is described and claimed in  
British Patent No. 2088877 (see Example 14 thereof).  
This compound has proven anti-inflammatory activity and  
is particularly useful for the treatment of respiratory  
30 disorders, particularly asthma. Fluticasone propionate  
has been obtained in a crystalline form, designated Form  
1, by dissolving the crude product (obtained, e.g. as  
described in British Patent No. 2088877) in ethyl  
acetate and then recrystallising. Standard spray-drying  
35 techniques have also been shown to lead only to the  
known Form 1 of fluticasone propionate. According to  
the present invention, fluticasone propionate may be

prepared in a new polymorphic form, designated Form 2. Form 2 may be characterised for example by its X-ray powder diffraction (XRPD) pattern (see *infra*).

5 The particulate products of the present invention are produced according to a supercritical fluid technique which we have developed.

10 The use of supercritical fluids (SCFs) and the properties thereof have been extensively documented, see for instance, J.W. Tom and P.G. Debendetti, "Particle Formation with Supercritical Fluids - A Review", *J. Aerosol. Sci.*, 22 (5), 555-584 (1991). Briefly, a supercritical fluid can be defined as a fluid at or 15 above its critical pressure ( $P_c$ ) and critical temperature ( $T_c$ ) simultaneously. Supercritical fluids have been of considerable interest, not least because of their unique properties. These characteristics include:

20 • High diffusivity, low viscosity and low surface tension compared with liquids.

25 • Large compressibility of supercritical fluids compared with the ideal gas implies large changes in fluid density for slight changes in pressure, which in turn results in highly controllable solvation power. Supercritical fluid densities typically range from 0.1- 0.9 g/ml under normal working conditions. Thus, 30 selective extraction with one supercritical fluid is possible.

35 • Many supercritical fluids are normally gases under ambient conditions, which eliminates the evaporation/concentration step needed in conventional liquid extraction.

• Most of the commonly used supercritical fluids create

non-oxidising or non-degrading atmospheres for sensitive and thermolabile compounds, due to their inertness and moderate temperatures used in routine working conditions. Carbon dioxide is the most extensively used 5 SCF due to its cheapness, non-toxicity, non-flammability and low critical temperature.

These characteristics have led to the development of several techniques of extraction and particle formation 10 utilising supercritical fluids. In particular, two particle formation methods have been identified:

Rapid expansion of supercritical solution (RESS) (see, for instance, J.W. Tom and P.G. Debenedetti, *supra*) 15 involves the dissolution of the solute of interest in a supercritical fluid, followed by rapid expansion of the resulting supercritical solution to atmospheric pressure, resulting in the precipitation of solute particles.

Gas Anti Solvent (GAS) recrystallisation (P.M. Gallagher et al, *Supercritical Fluid Science and Technology*, ACS Symp. Ser. 406, 134 (1989)) is particularly useful in 20 situations when the solvent of interest does not dissolve in, or has a very low solubility in, a supercritical fluid or a modified supercritical fluid. In this technique, the solute is dissolved in a 25 conventional solvent. A supercritical fluid such as carbon dioxide is introduced into the solution, leading to a rapid expansion of its volume. As a result, the 30 solvent power decreases dramatically over a short period of time, triggering the precipitation of the particles.

There is a need for techniques whereby a product may be 35 obtained with consistent and controlled physical criteria, including control of particle size and shape, quality of the crystalline phase, chemical purity and

enhanced handling and fluidising properties.

In addition, it would be advantageous to prepare micron-sized particles directly without the need to mill 5 products to this size range. Such milling leads to associated problems such as increased static charge and enhanced particle cohesiveness, as well as reduced yield of product. It also leads to highly stressed particles, which stress may affect the particles' dissolution after 10 administration.

Described in WO95/01324 is an apparatus for the formation of a particulate product in a controlled manner utilising a supercritical fluid particle 15 formation system. The disclosure of WO95/01324 is incorporated herein by this reference. The apparatus comprises a particle formation vessel with means for controlling the temperature of said vessel and means for controlling the pressure of said vessel, together with a 20 means for the co-introduction into said vessel of a supercritical fluid and a vehicle containing at least one substance in solution or suspension, such that dispersion and extraction of the vehicle occur substantially simultaneously by the action of the 25 supercritical fluid.

As used herein, the term "supercritical fluid" means a fluid at or above its critical pressure ( $P_c$ ) and critical temperature ( $T_c$ ) simultaneously. In practice, the 30 pressure of the fluid is likely to be in the range  $1.01 P_c - 7.0 P_c$ , and its temperature in the range  $1.01 T_c - 4.0 T_c$ .

The term "vehicle" means a fluid which dissolves a solid 35 or solids, to form a solution, or which forms a suspension of a solid or solids which do not dissolve or have a low solubility in the fluid. The vehicle can be

composed of one or more fluids.

As used herein, the term "supercritical solution" means a supercritical fluid which has extracted and dissolved  
5 the vehicle.

The term "dispersion" means the formation of droplets of the vehicle containing at least one substance in solution or suspension.

10 The term "particulate product" includes products in a single-component or multi-component (e.g. intimate mixtures of one component in a matrix of another) form.

15 It will be appreciated that, where necessary, the apparatus may additionally comprise a means for the collection of the particulate product, for example, a means for the retention of the product in the particle formation vessel, such as a filter, thus reducing loss  
20 of the product together with the resultant supercritical solution. An alternative means may involve a cyclone separating device.

25 The apparatus mentioned above and its use provide the opportunity for manufacturing dry particulate products with controlled particle size and shape by offering control over the working conditions, especially the pressure, utilising, for example, an automated back-pressure regulator such as model number 880-81 produced  
30 by Jasco Inc. Such an improved control eliminates pressure fluctuation across the particle formation vessel and ensures a more uniform dispersion of the vehicle (containing at least one substance in solution or suspension) by the supercritical fluid with narrow  
35 droplet size distribution during the particle formation process. There is little or no chance that the dispersed droplets will reunite to form larger droplets

since the dispersion occurs by the action of the supercritical fluid which also ensures thorough mixing with the vehicle and rapidly removes the vehicle from the substance(s) of interest, leading to particle 5 formation.

The simultaneous co-introduction of the vehicle containing at least one substance in solution or suspension and the supercritical fluid, according to the 10 method described herein, allows a high degree of control of parameters such as temperature, pressure and flow rate, of both vehicle fluid and supercritical fluid, at the exact point when they come into contact with one another.

15 Further advantages for particles formed as described herein include control over the quality of the crystalline and polymorphic phases, since the particles will experience the same stable conditions of 20 temperature and pressure when formed, as well as the potential of enhanced purity. This latter feature can be attributed to the high selectivity of supercritical fluids under different working conditions, enabling the extraction of one or more of the impurities from the 25 vehicle containing the substance of interest.

The means for the co-introduction of the supercritical fluid and the vehicle into the particle formation vessel preferably allows for them to be introduced with 30 concurrent directions of flow, and more preferably takes the form of a coaxial nozzle as described below. This ensures no contact between the formed particles and the vehicle fluid around the nozzle tip area. Such contact would reduce control of the final product size and 35 shape. Extra control over the droplet size, in addition to that provided by nozzle design, is achieved by controlling the flow rates of the supercritical fluid

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and the vehicle fluid. At the same time, retaining the particles in the particles formation vessel eliminates the potential of contact with the vehicle fluid that might otherwise take place on depressurising the 5 supercritical solution. Such contact would affect the shape and size, and potentially the yield, of the product.

Thus, in the apparatus described herein and in 10 WO95/01324 the means for the co-introduction of the supercritical fluid and the vehicle (containing at least one substance in solution or suspension) into the particle formation vessel preferably comprises a nozzle the outlet end of which communicates with the interior 15 of the vessel, the nozzle having coaxial passages which terminate adjacent to one another at the outlet end, at least one of the passages serving to carry a flow of the supercritical fluid, and at least one of the passages serving to carry a flow of the vehicle in which a 20 substance is dissolved or suspended.

Preferably, the opening at the outlet end (tip) of the nozzle will have a diameter in the range of 0.05 to 25 2 mm, more preferably between 0.1 and 0.3 mm, typically about 0.2 mm. The angle of taper of the outlet end will depend on the desired velocity of the fluids introduced through the nozzle; an increase in the angle may be used, for instance, to increase the velocity of the supercritical fluid introduced through the nozzle and 30 hence to increase the amount of physical contact between the supercritical fluid and the vehicle. Typically (although not necessarily) the angle of taper will be in the range of about 10° to about 50°, preferably between about 20° and about 40°, more preferably about 30°. The 35 nozzle may be made of any appropriate material, for example stainless steel.

In one embodiment, the nozzle has two coaxial passages, an inner and an outer. In another, preferred, embodiment, the nozzle has three coaxial passages, an inner, an intermediate and an outer. This latter design 5 allows greater versatility in use of the apparatus, since if necessary two vehicles may be introduced into the particle formation vessel with the supercritical fluid. Improved dispersion and finer particles can also be obtained if such a nozzle is used to introduce a flow 10 of the vehicle sandwiched between an inner and an outer flow of the supercritical fluid, since this ensures that both sides of the vehicle are exposed to the supercritical fluid. It is, however, to be appreciated that the nozzle may have any appropriate number of 15 coaxial passages.

The internal diameters of the coaxial passages may be chosen as appropriate for any particular use of the apparatus. Typically, the ratio of the internal 20 diameters of the outer and the inner passages may be in the range of from 2 to 5, preferably between about 3 and 5. Where an intermediate passage is included, the ratio of the internal diameters of the outer and intermediate passages may be in the range of from 1 to 3, preferably 25 between about 1.4 and 1.8.

Particular examples of such coaxial nozzles and their typical dimensions are illustrated in Figures 2A, 2B and 4 herein.

30 The temperature of the particle formation vessel may be maintained (preferably  $\pm 0.1^\circ\text{C}$ ) by means of a heating jacket or, more preferably, an oven. The pressure of the particle formation vessel is conveniently maintained 35 (preferably  $\pm 2$  bar) by means of a back-pressure regulator. It will be appreciated that such apparatus will be readily available from, for example,

manufacturers of supercritical fluid extraction equipment, for instance, from Jasco Inc., Japan.

The invention provides a method for the formation of a particulate fluticasone propionate product which comprises the co-introduction of a supercritical fluid and a vehicle containing at least fluticasone propionate in solution or suspension into a particle formation vessel, the temperature and pressure in which are controlled, such that dispersion and extraction of the vehicle occur substantially simultaneously by the action of the supercritical fluid.

Dispersion and extraction will also typically occur substantially immediately on introduction of the fluids into the particle formation vessel. Co-introduction of the supercritical fluid and the vehicle containing at least fluticasone propionate in solution or suspension preferably is effected using a nozzle of coaxial design.

Suitably the particle formation vessel used is as described in WO95/01324.

Suitable chemicals for use as supercritical fluids include carbon dioxide, nitrous oxide, sulphur hexafluoride, xenon, ethylene, chlorotrifluoromethane, ethane and trifluoromethane. Particularly preferred is carbon dioxide.

The supercritical fluid may optionally contain one or more modifiers, for example, but not limited to, methanol, ethanol, ethyl acetate, acetone, acetonitrile or any mixture thereof. When used, the modifier preferably constitutes not more than 20%, and more particularly constitutes between 1 and 10%, of the supercritical fluid.

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The term "modifier" is well known to those persons skilled in the art. A modifier (or co-solvent) may be described as a chemical which, when added to a supercritical fluid, changes the intrinsic properties of 5 the supercritical fluid in or around the critical point.

Regarding the choice of vehicle for the fluticasone propionate, where the fluticasone propionate is to be handled as a solution it should be soluble in the chosen 10 vehicle, and the chosen vehicle should be soluble in the chosen supercritical fluid. The choice of a suitable combination of supercritical fluid, modifier (where desired) and vehicle for any desired product will be well within the capabilities of a person of ordinary 15 skill in the art.

Suitable solvents may be, for example, methanol, ethanol, ethyl acetate, acetone, acetonitrile or any mixture thereof.

20 Control of parameters such as size, shape and crystal habit in the particulate product will be dependent upon the operating conditions used when carrying out the method of the invention. Variables include the flow 25 rates of the supercritical fluid and/or the vehicle containing substance(s), the vehicle used to dissolve the substance(s), the concentration of the substance(s) in the vehicle, and the temperature and pressure inside the particle formation vessel.

30 It will also be appreciated that the precise conditions of operation of the present apparatus will be dependent upon the choice of supercritical fluid and whether or not modifiers are present.

35 Table 1 lists the critical pressure and temperatures for some selected fluids:

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Table 1

| Fluid                  | P <sub>c</sub> (bar) | T <sub>c</sub> (°C) |
|------------------------|----------------------|---------------------|
| carbon dioxide         | 74                   | 31                  |
| nitrous oxide          | 72                   | 36                  |
| sulphur hexafluoride   | 37                   | 45                  |
| xenon                  | 58                   | 16                  |
| ethylene               | 51                   | 10                  |
| chlorotrifluoromethane | 39                   | 29                  |
| ethane                 | 48                   | 32                  |
| trifluoromethane       | 47                   | 26                  |

5 In practice, it may be preferable to maintain the  
 15 pressure inside the particle formation vessel  
 substantially in excess of the P<sub>c</sub> (for instance, 100-300  
 bar for carbon dioxide) whilst the temperature is above  
 the T<sub>c</sub> (e.g. 35-75°C for carbon dioxide).

20 The flow rates of the supercritical fluid and/or the  
 vehicle may also be controlled so as to achieve a  
 desired particle size, shape and/or form. Although the  
 flow ratio will be dependent on the desired  
 25 characteristics of the fluticasone propionate, typically  
 the ratio of the vehicle flow rate to the supercritical  
 fluid flow rate will be between 0.001 and 0.1,  
 preferably between 0.01 and 0.07, more preferably around  
 0.03.

30 The method described herein preferably additionally  
 involves collecting the particulate product following  
 its formation. It may also involve recovering the  
 supercritical solution formed, separating the components  
 of the solution and recycling one or more of those  
 35 components for future use.

According to a preferred aspect of the present invention, there is provided the compound fluticasone propionate in an easily handled and easily fluidised crystalline form, with a controlled particle size and 5 shape, and optionally also with a controlled morphology and level of agglomeration.

The invention also provides a new crystalline form of fluticasone propionate, designated Form 2 fluticasone propionate as described herein. The precise conditions 10 under which Form 2 fluticasone propionate is formed may be empirically determined; herein (see Example 5) we give a number of examples of methods which have been found to be suitable in practice.

15 As mentioned above, control of parameters such as size, shape and crystal habit in the particulate product will be dependent upon the operating conditions used when carrying out the method of the invention. By 20 appropriate adjustment of the variables of the process, the relative amounts of Forms 1 and 2 of fluticasone propionate produced with the apparatus described herein may be altered by the person skilled in the art. The experimental domains for each polymorphic form can be 25 determined empirically for the particular apparatus employed.

Conventionally crystallised fluticasone propionate, even 30 after micronisation (fluid milling), exists in a form with very poor flow characteristics, for example it is cohesive and statically charged, which results in difficulties in handling the drug substance in pharmaceutical formulation processes.

35 In another aspect of the present invention, there is provided fluticasone propionate in a form with a dynamic bulk density of less than 0.2 g cm<sup>-3</sup>. In a preferred

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aspect of the present invention, there is provided fluticasone propionate in a form with a dynamic bulk density in the range between 0.05 and 0.17 g cm<sup>-3</sup>, and, in particular, in the range between 0.05 and 0.08 g cm<sup>-3</sup>.

5

The dynamic bulk density (W) is indicative of a substance's fluidisability and is defined as:

10

$$W = \frac{(P-A)C}{100} + A$$

15

where P is the packed bulk density (g cm<sup>-3</sup>), A is the aerated bulk density (g cm<sup>-3</sup>) and C is the compressibility (%) where C is calculated by the equation:

20

$$C = \frac{P-A}{P} \times 10$$

Clearly, a low figure for W corresponds to a high degree of fluidisability.

25

When compared against conventionally crystallised fluticasone propionate, both before and after micronisation, fluticasone propionate of the present invention exhibits a significantly lower dynamic bulk density than the conventionally crystallised fluticasone propionate as illustrated in Table 7 (see Example 6 below).

35

It will be appreciated that in the case of an inhaled pharmaceutical, such as fluticasone propionate, it is particularly desirable to produce a drug substance which is readily fluidisable, thereby potentially improving its inhalation properties.

40

The fluticasone propionate of the present invention is observed to have improved handling and fluidising

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characteristics compared with conventionally crystallised fluticasone propionate.

5 Furthermore, the particle size and shape of provided fluticasone propionate of the present invention can be controlled as illustrated by the electronmicrographs herein.

10 Preferably, the fluticasone propionate of the present invention is within the particle size range suitable for pharmaceutical dosage forms to be delivered by inhalation or insufflation. A suitable particle size range for this use is 1 to 10 microns, preferably 1 to 5 microns. Particles generally have a uniform particle 15 size distribution, as measured by a uniformity coefficient of from 1 to 100, typically 1 to 20 e.g. 5 to 20.

20 The particle size distribution of the fluticasone propionate according to the invention may be measured by conventional techniques, for example by laser diffraction, by the "Twin Impinger" analytical process or by the "Cascade Impaction" analytical process. As used herein reference to the "Twin Impinger" assay means 25 "Preparations for Inhalation: Aerodynamic assessment of fine particles using apparatus A" as defined in the British Pharmacopoeia 1993, Addendum 1996, pages A522-527 as applied to a dry powder inhalation formulation. As used herein reference to the "Cascade Impaction" assay means "Preparations for Inhalation: Aerodynamic 30 assessment of fine particles using apparatus D" as defined in the British Pharmacopoeia 1993, Addendum 1996, page 527 as applied to a metered dose inhaler formulation. The preferred fluticasone propionate 35 according to the invention of mean particle size between 1 and 10 microns has been found to have a respirable fraction of 14% or more by weight.

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The fluticasone propionate of the present invention typically has a low cohesivity, for example of 0 to 20%, preferably 0 to 10% employing methods of measurement based on those described by R.L. Carr in Chemical 5 Engineering 1965, 163-168.

The fluticasone propionate according to the invention may be used to prepare a pharmaceutical composition which may be presented for use in a conventional manner 10 with the aid of a pharmaceutically acceptable carrier or excipient, optionally with supplementary medicinal agents. Preferred carriers include, for example, polymers e.g. starch and hydroxypropylcellulose, silicon dioxide, sorbitol, mannitol and lactose e.g. lactose 15 monohydrate. The compositions may be in a form suitable for administration by inhalation or insufflation, or for oral, buccal, parenteral, topical (including nasal) or rectal administration. Administration by inhalation or insufflation is preferred.

20 In a preferred pharmaceutical composition according to the invention the fluticasone propionate and carrier are co-crystallised together using the process and apparatus described herein to form multicomponent particles 25 comprising both fluticasone propionate and carrier. Such multicomponent particles represent a further aspect of the invention.

30 In a preferred aspect the invention provides a pharmaceutical composition in the form of a dry powder suitable for inhalation or insufflation which comprises fluticasone propionate according to the present invention and a suitable powder base such as lactose or starch, preferably lactose, as carrier. Especially 35 preferred are compositions comprising fluticasone propionate and lactose in the form of multicomponent particles. The dry powder composition may be presented

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in unit dosage form in, for example, capsules or cartridges of e.g. gelatin, or blister packs from which the powder may be administered with the aid of an inhaler or insufflator.

5

For administration by inhalation the fluticasone propionate made in accordance with the invention may be conveniently delivered in the form of an aerosol spray presentation from pressurised packs such as metered dose 10 inhalers, with the use of a suitable propellant, such as dichlorodifluoromethane or preferably a fluorocarbon or hydrogen-containing fluorocarbon such as HFA134a (1,1,1,2-tetrafluoroethane), HFA227 (1,1,1,2,3,3,3-heptafluoro-n-propane) or mixtures thereof. Such 15 aerosol spray presentations may include surfactants, e.g. oleic acid or lecithin; co-solvents, e.g. ethanol; or other excipients conventionally used in such formulations.

20 The formulations for administration by inhalation or insufflation are intended for administration on a prophylactic basis to humans suffering from allergic and/or inflammatory conditions of the nose, throat or lungs such as asthma and rhinitis, including hay fever.

25 Aerosol formulations are made so that each metered dose or "puff" of aerosol contains from 20 to 1000 micrograms, preferably 25 to 150 micrograms of fluticasone propionate of the invention. Administration may be several times daily, for example 2, 3, 4 or 8 30 times, giving for example 1, 2 or 3 doses each time. The overall daily dose with an aerosol will be within the range 100 micrograms to 10 mg, preferably 100 micrograms to 1.5 mg.

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There follows a brief description of the Figures:

Figure 1 shows a schematic design of an apparatus described herein.

5

Figure 2A shows a cross-section of a coaxial nozzle for use in the apparatus described herein.

10

Figure 2B shows a longitudinal section of a tip of a coaxial nozzle for use in the apparatus described herein.

Figures 3A and 3B show schematic designs of alternative apparatuses.

15

Figure 4 shows a longitudinal section of the tip of an alternative coaxial nozzle.

20

Figures 5 to 7 are scanning electron microscopy (SEM) photographs of fluticasone propionate, as prepared in Example 2.

Figure 8 is an (SEM) photograph of fluticasone propionate, as prepared in Example 3.

25

Figure 9 is an X-ray powder diffraction (XRPD) pattern of fluticasone propionate, as prepared in Example 2.

30

Figure 10 is an X-ray powder diffraction (XRPD) pattern of fluticasone propionate, as prepared in Example 3.

Figure 11 is a differential scanning calorimetry (DSC) profile of fluticasone propionate, as prepared in Example 2.

35

Figure 12 is a differential scanning calorimetry (DSC) profile of fluticasone propionate, as prepared in

## Example 3.

Figure 13 is a fourier transform infra-red (FTIR) spectrum of fluticasone propionate, as prepared in  
5 Example 2.

Figure 14 is a fourier transform infra-red (FTIR) spectrum of fluticasone propionate, as prepared in  
Example 3.

10 Figures 15 to 19 are HPLC chromatograms for the fluticasone propionate products as described in Example 4.

15 Figures 20 to 24 are XRPD patterns for the fluticasone propionate products as described in Example 5.

20 There follows a detailed description of a preferred embodiment of the apparatus and method described herein with reference to Figures 1, 2, 3 and 4. Figures 1 and 3 are simplified diagrammatic flow sheets of apparatus and Figures 2A, 2B and 4 show nozzles which may be used therein.

25 Referring to Figure 1, the apparatus includes a particle formation vessel 6. This is typically a standard reaction vessel, for instance of the type available from Keystone Scientific Inc., of an appropriate capacity for the particular use to which it is to be put. The  
30 temperature and pressure of the vessel are maintained at constant desired level, by means of an oven 7 and back-pressure regulator 8, respectively.

35 In use, the system is initially pressurised and stable working conditions are met. A suitable gas, for example, carbon dioxide, is fed from source 1 via conduit 11 to a cooler 2, to ensure liquification, and

is fed by conduit 12 to a pump 4. From there it is fed by conduit 13 to the vessel 6 via a nozzle 20. A solution or dispersion of a solid of interest, in this case fluticasone propionate, in a suitable vehicle, for 5 example methanol, is drawn from source 5 by a conduit 14 to a pump 3 and is fed by conduit 15 to the vessel 6 via a nozzle 20.

The nozzle 20 may be as shown in either Figure 2 (A and 10 B) or Figure 4. That shown in Figure 2 comprises coaxial inner and outer tubes 30 and 40, respectively. These define an inner passage 31 and an outer passage 41. The tubes 30 and 40 have conically tapering end portions 32 and 42, respectively. The tips of the end 15 portions 32 and 42 define respective orifices 33 and 43, with the orifice 43 being a short distance downstream of the orifice 33. As indicated in Figure 2B, the angle of taper of the end portion 42 is about 30° in this (non-limiting) example.

20 The alternative nozzle illustrated in Figure 4 comprises three coaxial tubes 50, 60 and 70 which define an inner passage 51, an intermediate passage 61, and an outer passage 71 respectively. Tubes 60 and 70 have conically tapering end portions 62 and 72, the angle of taper of 25 the end portion 72 being about 30° in this example.

The nozzle of Figure 4 allows three fluids to be introduced into the vessel 6 at the same time, leading 30 to greater versatility in use of the apparatus. For instance, it is possible to add through one of the three passages a desired carrier or other additive intended to form part of, or be mixed with, the final particulate product. The additive is then dispersed simultaneously 35 with the substance of primary interest. Also, in situ reactions may be carried out immediately prior to dispersion by the supercritical fluid, by introducing

two or more reactants in two separate vehicles through two of the nozzle passages, the reaction occurring at the passage outlets either immediately prior to, or on, dispersion.

5

Alternatively, the nozzle of Figure 4 may be used to introduce a flow of the vehicle (passage 61) sandwiched between an inner and an outer flow of the supercritical fluid (passages 51 and 71). This leads to improved 10 dispersion of the vehicle, and hence to greater control over, and uniformity of, particle size in the final product; indeed it makes possible the formation of finer products than may be achieved using a two-passage nozzle.

15

In the nozzle shown, inner tube 50 has an internal diameter of 0.25 mm; intermediate tube 60 has an internal diameter of 0.53 mm; and outer tube 70 has an internal diameter of 0.8 mm and an outside diameter of 20 1.5 mm. The tip opening (73) has an internal diameter of 0.2 mm. The tubes are all made of stainless steel.

However, the nozzle may be made of any appropriate material and have any suitable dimensions. For 25 instance, the internal diameters may be in the ranges 0.05-0.35 mm (inner); 0.25-0.65 mm (intermediate); and 0.65-0.95 mm (outer), preferably between 0.1 and 0.3 mm (inner); 0.3 and 0.6 mm (intermediate); and 0.07 and 0.9 mm (outer). The tip opening is likely to have an 30 internal diameter in the range of 0.1-0.3 mm, preferably between 0.18 and 0.25 mm.

In the apparatus of Figure 1, the supercritical fluid is fed under pressure (at high flow rate when compared with 35 the flow rate of the vehicle) through for example the inner nozzle passage 31 of the nozzle shown in Figure 2, and the solution or suspension of the solid of interest

in a vehicle (hereinafter referred to as the "liquid") is simultaneously fed under pressure through the outer passage 41. It is believed that the high velocity supercritical fluid emerging from the orifice 33 causes 5 the liquid emerging from the end of outer passage 41 to be broken up into droplets from which the vehicle is substantially simultaneously extracted by the supercritical fluid to result in the formation of particles of the solid previously held in the vehicle. 10 It is to be understood, however, that although it is believed that this is what occurs, we do not wish to be bound by this theoretical explanation, and the actual physical processes occurring may not be precisely as just indicated.

15 Also, although the configuration has been described in which the supercritical fluid passes through the inner passage 31 and the vehicle passes through the outer passage 41, the configuration may be reversed, with the 20 supercritical fluid in the outer passage 41 and the vehicle in the inner passage 31. Similarly in the nozzle of Figure 4, any one of the three passages may be used to carry any one of a number of desired fluids, as appropriate.

25 The nozzle 20 ensures dispersion of the vehicle containing the solid of interest, the shearing action of the high velocity supercritical fluid and also thorough mixing of the dispersed vehicle with the supercritical 30 fluid which simultaneously extracts the vehicle from the dispersed liquid, resulting in substantially immediate particle formation of the solid of interest. Because the supercritical fluid and vehicle are introduced coaxially, and dispersion occurs substantially 35 simultaneously with vehicle extraction, a very high degree of control is possible of the conditions (e.g. pressure, temperature and flow rate) affecting particle

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formation, at the exact time when it occurs.

The particles formed are retained in the particle formation vessel by collecting means 21. The resultant 5 supercritical solution is fed by conduit 16 to a back-pressure regulator 8 and is then fed by conduit 17 to a separation vessel 9 where it expands to cause the supercritical fluid to separate as a gas from the liquid vehicle. The gas may be fed by conduit 18 to a tank 10 and returned by conduit 19 to the cooler 2. The vehicle 10 may also be collected for subsequent use. Means, not shown, may be provided to smooth the flow pulse of fluids and vehicles produced by pumps 3 and 4, so as to eliminate, or at least reduce, any flow pulsations.

15

When sufficient particle formation has occurred in the vessel 6, it is flushed through with clean, dry supercritical fluid, so as to ensure removal of any residual vehicle. The vessel can then be depressurised 20 and particulate product removed.

The alternative apparatuses shown schematically in Figures 3A and 3B are for use in continuous particle formation. That shown in Figure 3A includes two 25 particle formation vessels 6a and 6b, each of the type shown in Figure 1 and each including an inlet nozzle 20 and a particle collecting means (such as a filter) 21. Oven 7 serves both vessels.

30 In the apparatus in Figure 3A, valve A controls the supply of the supercritical fluid and the vehicle (containing the substance of interest) to the two vessels 6a and 6b, and one-way valves E and F control the outlets from the two vessels to the back-pressure 35 regulator 8. Valve D controls the supply of the vehicle to valve A. Valves B and C are needle valves, and items 80 and 81 are vents.

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The apparatus may be "continuously" operated as follows. Valve A is firstly set to supply fluids to vessel 6a, in which particle formation is allowed to occur, as described in connection with Figure 1. Valve E is set 5 so that the resultant supercritical solution may drain from vessel 6a to the back-pressure regulator 8 for subsequent recycling.

When sufficient particle formation has occurred, valve D 10 is closed to stop the flow of vehicle, whilst the supercritical fluid continues to flow through vessel 6a to dry (flush) the products. Valve A is then set to supply fluids to the empty vessel 6b and valve D re-opened, whilst valve B is opened so as slowly to 15 depressurise vessel 6a. One-way valve E eliminates any chance of a back-flow from vessel 6b or of disruption of the particle formation process now occurring in vessel 6b. Vessel 6a is removed for collection of the product, and then refitted and repressurised ready for re-use. 20 Supercritical solution drains from vessel 6b via valve F, which is set appropriately.

Once particle formation in vessel 6b is complete, the valves are set back to allow it to continue in vessel 25 6a, whilst 6b is flushed and emptied. In this way, particle formation in the apparatus can continue uninterrupted.

The apparatus shown in Figure 3B includes only one 30 particle formation vessel 6, which does not contain any particle collecting means, and two particle collection vessels 25a and 25b downstream of vessel 6. The supercritical fluid carries the formed particles to the collection vessels 25a and 25b.

35

The apparatus also includes an inlet nozzle 20, two vents 26, a back pressure regulator 27, an oven 7 and

valves A - H. Supercritical fluid and solution (vehicle) are fed to the nozzle 20 where shown.

The apparatus might be used as follows. Initially, 5 (valves C, D, E, and F closed) the system is pressurised and stable working conditions are met; valves B and H are then closed, driving the flow of supercritical fluid through valve A only. The vehicle and substance of interest are introduced into vessel 6 and the particles 10 formed are transported by the supercritical fluid via valve A to collection vessel 25a which contains a particle retention device. The retention device is placed at the outlet of the vessel to ensure maximum collection volume. The solid-free supercritical 15 solution (the supercritical fluid and the vehicle) flows across valve G to the back pressure regulator 27. On emerging from the back pressure regulator the supercritical solution expands into a large pressure 20 resistant vessel (not shown), where the vehicle separates from the gas and both can be recycled.

When the collection vessel 25a is full, switching takes place, closing valves A and G and simultaneously opening valves B and H. This allows the flow of the 25 supercritical solution, emerging from vessel 6, into the second collection vessel 25b. Valves C and G are opened after flow switching to ensure a high flow of supercritical fluid to flush the full collection vessel 25a, i.e. the supercritical solution volume is replaced 30 by a supercritical fluid volume. It is estimated that 1-2 times the volume of the collection vessel, of the supercritical fluid, ensures a dry powder. The flushing time is generally short owing to the fact that the particles are occupying the volume of the collection 35 vessel. After flushing, valves C and G are closed and valve F (a needle valve) is slowly opened to the full collection vessel 25a. Since the particulate product

takes up the vessel volume only a small amount of supercritical fluid is discharged, mainly the internal volume of the fittings involved.

5 The full collection vessel 25a is removed and the dry powder collected. After refitting and repressurising via valve C, the vessel is ready for re-use as soon as the second collection vessel 25b, which has meantime been collecting product from vessel 6, is full.

10 10 The benefits of using the apparatus of Figure 3B include:

15 15 1. The elimination of depressurising and pressurising steps of the reaction vessel every time product is collected. This could mean considerable reductions in the amounts of fluids being discharged, in particular when using a large volume particle formation vessel (scaling up) or expensive high purity gases.

20 20 2. Significant time saving during the flushing (drying) procedure. In a batch particle formation process only a rather small volume of the reaction vessel is occupied by the product and the remaining volume (where dispersion takes place) is taken up by the supercritical solution. This mixture will eventually be replaced by at least the same volume of the supercritical fluid in the flushing procedure, which can therefore take a long time when scaled up.

25 25 3. The environment and workers are less exposed to the products during the recovery step. In some cases it is difficult to collect products directly from a large reaction vessel due to handling inconvenience or because the products of interest are sensitive to

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light, oxygen or humidity, which might affect their characteristics or purity.

The invention is further illustrated by the following  
5 non-limiting examples. Examples 1 to 9, illustrating  
the preparation of fluticasone propionate and its  
physical properties, were carried out using apparatus of  
the type illustrated in Figures 1 and 2, using a 32 ml  
particle formation vessel and a two-passage coaxial  
10 nozzle having the following dimensions:

|             | outer diameter | inner diameter |
|-------------|----------------|----------------|
| outer tube: | 1.58 mm        | 0.75 mm        |
| inner tube: | 0.63 mm        | 0.20 mm        |

15 The tip orifice (43 in Figure 2B) was 0.32 mm in diameter, and both the inner and outer tubes were made of stainless steel.

20 Example 1  
Particle Size Distribution

25 The data for four samples of fluticasone propionate of the present invention, produced using the method and apparatus described herein, are presented in Table 2 below. The particle size was determined by laser diffraction (Malvern Mastersizer).

30 Sample 1 was produced using a solution of fluticasone propionate in acetone (2.5% w/v) which was co-introduced with CO<sub>2</sub> at 300 bar, 35°C and a flow rate ratio of 0.014 via a coaxial nozzle into the particle formation vessel.

35 Sample 2 was produced using a solution of fluticasone propionate in acetone (0.5% w/v) which was co-introduced with CO<sub>2</sub> at 100 bar, 35°C and a flow rate ratio of 0.014 via a coaxial nozzle into the particle formation vessel.

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Sample 3 was produced using a solution of fluticasone propionate in ethyl acetate (0.5% w/v) which was co-introduced with CO<sub>2</sub> at 100 bar, 75°C and a flow rate ratio of 0.043 via a coaxial nozzle into the particle formation vessel.

Sample 4 was produced using a solution of fluticasone propionate in acetone (2.5% w/v) which was co-introduced with CO<sub>2</sub> at 100 bar, 75°C and a flow rate ratio of 0.014 via a 3-component coaxial nozzle into the particle formation vessel.

The data presented in Table 2 show that the particle size can be manipulated depending on the conditions. The particle size data for sample 4 indicate that a particle size similar to that of conventionally crystallised fluticasone propionate (micronised) can be achieved. The uniformity index is not significantly different from that of the conventionally crystallised fluticasone propionate (micronised).

Table 2

|    | Mean particle size ( $\mu\text{m}$ )                            | % $< 5\mu\text{m}$ | % $< 10\mu\text{m}$ | Uniformity Index |
|----|---|--------------------|---------------------|------------------|
| 25 | Conventionally crystallised fluticasone propionate (micronised) | 1-3                | Typically > 90      | Typically > 95   |
| 30 | Sample 1  | 5.2                | 48                  | 81               |
|    | Sample 2  | 10.4               | 20                  | 48               |
|    | Sample 3  | 31.0               | 6                   | 13               |
|    | Sample 4  | 2.8                | 68                  | 83               |

Example 2Particle Shape Data

5

The particle shape was determined by scanning electron microscopy. Data for three samples of fluticasone propionate of the present invention, produced using the method and apparatus described herein, are presented in Figures 5 to 7.

10

Sample 5 was produced using a solution of fluticasone propionate in methanol (0.5% w/v) which was co-introduced with CO<sub>2</sub> at 100 bar, 75°C and a flow rate ratio of 0.043 via a coaxial nozzle into the particle formation vessel. The particle shape is described as acicular with a high aspect ratio up to 200:1 (Fig.5).

15

Sample 6 was produced using a solution of fluticasone propionate in acetone (1.5% w/v) which was co-introduced with CO<sub>2</sub> at 200 bar, 55°C and a flow rate ratio of 0.029 via a coaxial nozzle into the particle formation vessel. The particle shape is described as flake-like (Fig.6).

20

Sample 7 was produced using a solution of fluticasone propionate in acetone (2.5% w/v) which was co-introduced with CO<sub>2</sub> at 100 bar, 75°C and a flow rate ratio of 0.014 via a coaxial nozzle into the particle formation vessel. The particle shape is described as equant (Fig.7).

30

Example 3Reproducibility

35

Three different solutions of fluticasone propionate in acetone (1.5 %w/v) were co-introduced with CO<sub>2</sub> at 200 bar, 55°C and a flow rate ratio of 0.029 via a coaxial

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to the particle formation vessel on three different days (Samples 6, 8, 9). The size, particle shape, polymorphic form and impurity profile were examined.

5 The particle size, particle shape, polymorphic form and impurity profile data show that the technique is reproducible when the same crystallising parameters are used.

10 a) Particle Size

The particle size was determined by laser diffraction (Malvern Mastersizer). The data are shown below in Table 3.

15 Table 3

|             | Mean Particle size ( $\mu\text{m}$ ) | % $<5\mu\text{m}$ | % $<10\mu\text{m}$ | Uniformity Index |
|-------------|--------------------------------------|-------------------|--------------------|------------------|
| Sample 6    | 9.7                                  | 24                | 51                 | 10               |
| Sample 8    | 9.4                                  | 23                | 53                 | 11               |
| 20 Sample 9 | 9.6                                  | 23                | 52                 | 12               |

b) Particle Shape

25 The particle shape has been determined by scanning electron microscopy. Data for samples 6 and 8 are shown in Figures 6 and 8 respectively.

c) Polymorphic form

30 The polymorphic form has been determined by X-ray powder diffraction, differential scanning calorimetry (DSC) and fourier transform infra-red (FTIR) spectroscopy for samples 6 and 8. Data are shown in Figures 9 to 14.

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Figs. 9, 11 and 13 relate to sample 6, whilst Figs. 10, 12 and 14 relate to sample 8.

d) Impurity Profile

5 The impurity profile has been determined by HPLC for samples 6 and 8. The data are shown in Table 4 and Figures 16 and 17 (relating to samples 6 and 8 respectively).

10

Example 4

Impurity Profile

15 The impurity profile has been determined by HPLC for samples 5, 6, 8, 10 and compared to conventionally crystallised fluticasone propionate. The data are shown in Table 4 and Figures 15 to 19 (which relate to samples 5, 6, 8, 10 and conventionally crystallised fluticasone propionate respectively).

20

Sample 10 was produced using a solution of fluticasone propionate in acetone (0.5% w/v) which was co-introduced with CO<sub>2</sub> at 300 bar, 35°C and a flow rate ratio of 0.043 via a coaxial nozzle into the particle formation vessel.

25

30 The data show that fluticasone propionate produced by the present invention does not alter the impurity profile when compared to conventionally crystallised fluticasone propionate. However, there is a small reduction in the total impurities (% w/w) for fluticasone propionate produced by the present invention.

Table 4

|    |   | Total Impurities (% w/w) | Most Abundant Impurity (% w/w) | 2nd most Abundant Impurity (% w/w) | Number of Impurities |
|----|---|--------------------------|--------------------------------|------------------------------------|----------------------|
| 5  | Conventionally crystallised fluticasone propionate (micronised) | 0.60                     | 0.20                           | 0.14                               | 5                    |
|    | Sample 4  | 0.45                     | 0.12                           | 0.12                               | 5                    |
| 10 | Sample 6  | 0.51                     | 0.14                           | 0.12                               | 5                    |
|    | Sample 8  | 0.51                     | 0.14                           | 0.13                               | 5                    |
|    | Sample 10   | 0.59                     | 0.17                           | 0.14                               | 5                    |

(Note: Limit of quantification = 0.045% w/w)

15 Example 5

Crystallinity and Polymorphism

X-ray powder diffraction (XRPD) patterns were generated using either a Siemens D 5000 or Philips X'pert MPD. The powders were scanned over a 2θ angle range of either 1.5° to 60°, or 0° to 35°, with a 0.02-0.05° step at 3-15 seconds count time using CuKα radiation. The Siemens D5000 measures intensity as counts per second and the Philips X'pert MPD measures intensity as counts.

25 Data for Samples 2, 6 and 11 were compared to those of conventionally crystallised fluticasone propionate. (See Figures 20, 9, 21 and 22 respectively.)

30 The production of Samples 2 and 6 is described above. Sample 11 was produced using a solution of fluticasone propionate in acetone (0.5% w/v) which was co-introduced with CO<sub>2</sub> at 300 bar, 75°C and a flow rate ratio of 0.014

via a coaxial nozzle into the particle formation vessel.

The data show that the crystallinity can be controlled by the crystallisation parameters. Using the technique 5 presented in the patent, the crystallinity of the fluticasone propionate can be improved significantly over conventionally crystallised fluticasone propionate.

As mentioned above, the relative amounts of Forms 1 and 10 2 of fluticasone propionate produced with the apparatus described herein may be altered by appropriate adjustment of the variables of the process of the invention. The experimental domains for each polymorphic form can be determined empirically for the particular apparatus 15 employed. Using the preparative process and apparatus described herein, it was found that Samples 1, 2, 5-12 and 14-16 were Form 2 fluticasone propionate; Samples 3 and 17 were a mixture of Forms 1 and 2 fluticasone propionate; and Sample 13 is Form 1 fluticasone 20 propionate.

The two polymorphic forms of fluticasone propionate are well characterised by their XRPD traces. Table 5 shows 25 the key 2θ peaks for identification of the two polymorphic forms of the fluticasone propionate by XRPD. Figure 23 shows the XRPD traces of Forms 1 and 2 fluticasone propionate overlaid.

30

Table 5

| Polymorph | Primary Peaks (°) | Secondary Peaks (°) |      |      |      |      |      |      |  |
|-----------|-------------------|---------------------|------|------|------|------|------|------|--|
|           |                   | 11.5                | 12.4 | 13.1 | --   | 14.9 | --   | 15.8 |  |
| Form 1    | 7.9 10.0          |                     |      |      |      |      |      |      |  |
| Form 2    | 7.6 9.8           | --                  | --   | 13.0 | 13.6 | --   | 15.2 | --   |  |

Traces for the fluticasone propionate produced according to the present invention are qualitatively different from the trace for the conventionally crystallised fluticasone propionate. Conventionally crystallised fluticasone propionate ("Form 1") has been found to have a monoclinic crystal structure with

$$a = 7.722\text{\AA}, b = 14.176\text{\AA}, c = 11.290\text{\AA}, \beta = 98.458^\circ$$

10 In contrast, the XRPD traces of the fluticasone propionate produced according to the present invention ("Form 2") was analysed and proved to be a pure polymorph with an orthorhombic structure having

15  $a = 23.404\text{\AA}, b = 14.048\text{\AA}, c = 7.6953\text{\AA},$   
all angles  $90^\circ$

20 Form 2 was found to be stable to conversion to Form 1: after 62 weeks at ambient temperature and humidity, no conversion was seen. Figure 24 shows the stability of Form 2 fluticasone propionate by XRPD.

25 The water content of the two crystalline forms of fluticasone propionate produced with the apparatus described herein has also been determined, and compared with that of conventionally crystallised and micronised fluticasone propionate. The results are shown in Table 6.

Table 6

|    |   | Water content (% w/w) |
|----|---|-----------------------|
| 5  | Conventionally crystallised fluticasone propionate (micronised) | < 0.1                 |
|    | Form 1 fluticasone propionate prepared as described herein      | < 0.1                 |
| 10 | Form 2 fluticasone propionate prepared as described herein      | > 0.3                 |

Example 6Bulk Density

15

The dynamic bulk density for conventionally crystallised fluticasone propionate (micronised and non-micronised) and fluticasone propionate of the present invention are shown in Table 7.

20

Sample 12 was produced using a solution of fluticasone propionate in ethyl acetate (0.5% w/v) which was co-introduced with CO<sub>2</sub> at 300 bar, 35°C and a flow rate ratio of 0.043 via a coaxial nozzle into the particle formation vessel.

30

Sample 13 was produced using a solution of fluticasone propionate in acetonitrile (0.5% w/v) which was co-introduced with CO<sub>2</sub> at 100 bar, 75°C and a flow rate ratio of 0.043 via a coaxial nozzle into the particle formation vessel.

Table 7

|    |   | Dynamic Bulk Density (W)<br>g cm <sup>-3</sup> |
|----|---|--|
| 5  | Conventionally crystallised fluticasone propionate (micronised)     | 0.21   |
|    | Conventionally crystallised fluticasone propionate (non-micronised) | 0.21   |
| 10 | Sample 12   | 0.05   |
|    | Sample 13   | 0.06   |
|    | Sample 3  | 0.17   |

15 The dynamic bulk density of fluticasone propionate of the present invention is significantly lower than for conventionally crystallised fluticasone propionate (micronised and non-micronised).

20 The data presented in Table 7 show that dynamic bulk density of fluticasone propionate of the present invention can be controlled using the crystallisation parameters of the method described within the patent.

#### Example 7

##### Static Charge Test

25 The relative static charge of the fluticasone propionate of the present invention can be controlled by the crystallisation parameters. The data indicate there are no significant reductions in relative static of the fluticasone propionate of the present invention when compared to conventional micronised fluticasone propionate. Fluticasone propionate collected from the particle formation vessel using the apparatus described

is dry and easily handled. Conventional micronised fluticasone propionate is cohesive, difficult to handle and statically charged.

5 A simple test was devised to ascertain a relative static charge based on the quantity of drug remaining coated to the walls of a vial after rolling a predetermined quantity of drug in the vial for 2 minutes. The greater the amount of the drug remaining on the vial, the higher  
10 the relative static charge associated with the drug substance. The results are displayed in Table 8.

Table 8

| 15 | Sample   | % Drug retained<br>on vial |
|----|--|----------------------------|
|    | Conventionally crystallised<br>fluticasone propionate (micronised) | 4.5                        |
|    | Sample 14  | 3.8                        |
| 20 | Sample 15  | 4.2                        |
|    | Sample 16  | 7.6                        |
|    | Sample 17  | 8.6                        |

25 Sample 14 was produced using a solution of fluticasone propionate in acetone (2.5% w/v) which was co-introduced with CO<sub>2</sub> at 300 bar, 75°C and a flow rate ratio of 0.043 via a coaxial nozzle into the particle formation vessel.

30 Sample 15 was produced using a solution of fluticasone propionate in acetone (3.5% w/v) which was co-introduced with CO<sub>2</sub> at 90 bar, 85°C and a flow rate ratio of 0.028 via a coaxial nozzle into the particle formation vessel.

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5        Sample 16 was produced using a solution of fluticasone propionate in ethyl acetate (0.5% w/v) which was co-introduced with CO<sub>2</sub> at 300 bar, 35°C and a flow rate ratio of 0.043 via a coaxial nozzle into the particle formation vessel.

10        Sample 17 was produced using a solution of fluticasone propionate in acetone (0.5% w/v) which was co-introduced with CO<sub>2</sub> at 100 bar, 75°C and a flow rate ratio of 0.043 via a nozzle into the particle formation vessel.

Example 8

Twin Impinger Test

15        A small quantity of drug was filled into each blister of a 4-blister Rotadisk™. The contents of each Rotadisk™ were emptied, via a Diskhaler™, into the Twin Impinger apparatus set to an airflow rate of 60 litres per minute. Each stage of the Twin Impinger apparatus contained a quantity of dissolving solvent, methanol, (stage 1, 7 ml and stage 2, 30 ml). The Rotadisk™ and Diskhaler™ were washed with methanol and the resultant solution made up to 50 ml. The stage 1 of the Twin Impinger was washed with methanol and the resultant solution made up to 50 ml. The stage 2 of the Twin Impinger was washed with methanol and the resultant solution made up to 50 ml. The solutions were assayed by UV spectrophotometry and the quantity of drug delivered to each stage of the Twin Impinger apparatus calculated. The results are displayed in Table 9. Table 10 displays the size data of the samples used.

Table 9

| Drug Deposition as % of Total Drug Recovered |   |         |         |           |      |
|--|---|---------|---------|-----------|------|
|  | Device  | Stage 1 | Stage 2 | Delivered | Dose |
| 5  | Conventionally crystallised fluticasone propionate (micronised) | 40      | 43      | 17        | 60   |
| 10   | Sample 15   | 36      | 47      | 17        | 64   |
|  | Sample 16   | 28      | 58      | 14        | 72   |

15

Table 10

|    | Mean Particle size ( $\mu\text{m}$ ) | % $<5\mu\text{m}$ | % $<10\mu\text{m}$ | Uniformity Index |
|----|--------------------------------------|-------------------|--------------------|------------------|
| 20 | Sample 15                            | 4.0               | 61                 | 84               |
|    | Sample 16                            | 8.4               | 27                 | 59               |

25 The stage 2 deposition represents the fine particle mass (respirable dose) reaching the deep lung. The delivered dose (sum of stage 1 and stage 2) represents the total dose available for inhalation and the emptying efficiency of the drug from the device.

30 Fluticasone propionate of the present invention shows no significant improvement in stage 2 deposition. An interesting feature of the present invention is that the 35 supercritical fluid crystallised fluticasone propionate with a particle size greater than that of conventionally crystallised fluticasone propionate (micronised) gives

an equivalent deposition (respirable dose) in the stage 2 of the Twin Impinger.

Fluticasone propionate of the present invention shows an 5 improved delivered dose indicating the drug is emptied well from the device and is presenting a greater quantity of drug for inhalation. Again it is an interesting feature of the present invention that the supercritical fluid crystallised fluticasone propionate 10 with a particle size greater than that of conventionally crystallised fluticasone propionate (micronised) gives a greater delivered dose.

These data indicate that fluticasone propionate of the 15 present invention has improved fluidisability and flow properties.

Example 9

20 Solvent Content Test

The solvent content of fluticasone propionate of the present invention was investigated by Nuclear Magnetic Resonance (NMR) and compared to that of conventional 25 crystallised fluticasone propionate. Each sample was tested for acetone content. (The conventionally crystallised fluticasone propionate was crystallised from acetone.) In addition, sample 5 was crystallised from methanol and therefore tested for methanol. 30 Table 11 displays the solvent content data for each sample.

- 40 -

Table 11

|   |  | Acetone<br>Content<br>(% w/w) | Methanol<br>Content<br>(% w/w) |
|---|--|-------------------------------|--------------------------------|
| 5 | Conventionally crystallised<br>fluticasone propionate (micronised) | typically 0.7                 | N/A                            |
|   | Sample 5   | Not Detected                  | Not Detected                   |
|   | Sample 8   | Not Detected                  | N/A                            |
|   | Sample 15  | Not Detected                  | N/A                            |

10

The data indicate that there are no detectable levels of residual solvent within the fluticasone propionate of the present invention. Lack of residual solvent in the 15 samples is consistent with GAS (Gas Anti-Solvent) recrystallisation and RESS (Rapid Expansion of Supercritical Solutions).

20 The benefits of no residual solvent within the sample may include: improved stability due to lack of solvent-drug interactions on temperature and humidity storage; reduced crystal imperfections and improved crystal structure due to lack of solvent occlusions.

25

**Examples 10-12: Performance of Metered Dose Inhalers**

In the following tests, two types of metered dose inhaler (MDI) were manufactured, both containing 30 fluticasone propionate and HFA134a. Inhaler Type A was a 125 microgram, 120 actuation model. Inhaler Type B was a 50 microgram, 120 actuation model.

Inhalers Type A were prepared by dispensing 20 mg drug into an 8 ml Presspart aluminium can. The can was closed by crimping on a Valois DF60 63 microlitre Valve before pressure-filling the canister with 12 g of 5 Propellant HFA134a. Inhalers Type B were prepared in the same way but employing only 8 mg drug.

The performance of MDI's has been measured based on drug deposition on the can, valve and actuator; dose 10 delivered through use; and respirable dose.

The following crystallisation conditions were used in preparing samples for filling into the aforementioned MDI's.

15 Sample 18 was produced using a solution of fluticasone propionate in acetonitrile (2.0%w/v) which was co-introduced with CO<sub>2</sub> at 300 bar, 35°C and a flow rate ratio of 0.043 via a coaxial nozzle into the particle 20 formation vessel.

Sample 19 was produced using a solution of fluticasone propionate in acetone (2.5%w/v) which was co-introduced with CO<sub>2</sub> at 100 bar, 35°C and a flow rate ratio of 0.043 25 via a coaxial nozzle into the particle formation vessel.

Sample 20 was produced using a solution of fluticasone propionate in acetone (2.5%w/v) which was co-introduced with CO<sub>2</sub> at 100 bar, 35°C and a flow rate ratio of 0.014 30 via a coaxial nozzle into the particle formation vessel.

Sample 21 was produced using a solution of fluticasone propionate in acetone (2.5%w/v) which was co-introduced with CO<sub>2</sub> at 100 bar, 35°C and a flow rate ratio of 0.033 35 via a coaxial nozzle into the particle formation vessel.

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5        Sample 22 was produced using a solution of fluticasone propionate in acetonitrile (2.0%w/v) which was co-introduced with CO<sub>2</sub> at 100 bar, 35°C and a flow rate ratio of 0.022 via a coaxial nozzle into the particle formation vessel.

Example 10

Drug Deposition

10      The drug deposited on the can, valve and actuator were measured at the beginning of use of the inhaler (after actuations 1 & 2) and at the end of use of the inhaler (after actuations 119 & 120). After the appropriate number of actuations, the inhaler exterior is washed in acetonitrile to remove any residual drug deposited on the surface. The drug found on the actuator is washed into a suitable container with 50 ml acetonitrile/water (50:50 v/v). The inhaler is then frozen in liquid nitrogen, the valve removed quickly and the contents of the drug suspension emptied into a suitable container. The propellant from the suspension is allowed to evaporate and the drug remaining is dissolved in 50 ml acetonitrile/water (50:50 v/v). The drug on the valve components is washed into a suitable container with 50 ml acetonitrile/water (50:50 v/v). The drug on the can is also washed into a suitable container with 50 ml acetonitrile/water (50:50 v/v). The resultant solutions are assayed by HPLC.

20

25

30      Tables 12 and 13 present the drug deposition profile for Inhalers Types A and B respectively.

35      Fluticasone propionate of the present invention shows significantly lower drug deposition on the can, valve and actuator than conventionally produced fluticasone propionate. As a result of the lower drug deposition, the concentration of drug in the suspension is higher,

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leading to higher quantities of drug being delivered from the inhaler. This is confirmed in Examples 11 (Dose Through Use) and 12 (Cascade Impactor Testing), which show higher dose delivery through the life of the  
5 inhaler.

Table 12

10 Drug Deposition Profile for Inhaler Type A

|    | Drug   | Conventionally crystallised fluticasone propionate (micronised) | Fluticasone propionate of the present invention (sample 18) | Fluticasone propionate of the present invention (sample 20) |
|----|--|---|---|---|
| 15 | Amount of drug in Suspension (mg)                        | 17.7  | 18.1  | 18.9  |
| 20 | Can Deposition at the beginning of use (mg)              | 1.0   | 0.8   | 0.5   |
| 25 | Valve Deposition at the beginning of use (mg)            | 0.7   | 0.5   | 0.3   |
| 30 | Total Drug Deposition at the beginning of use (mg)       | 1.7   | 1.3   | 0.8   |
|    | Actuator Deposition at the beginning of use (micrograms) | 15  | 12  | 11  |

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Table 13

## Drug Deposition Profile for Inhaler Type B

| 5  | Drug   | Conventionally crystallised fluticasone propionate (micronised) | Fluticasone propionate of the present invention (sample 21) | Fluticasone propionate of the present invention (sample 22) |
|----|--|---|---|---|
| 10 | Amount of drug in Suspension (mg)                  | 6.4   | 7.1   | 7.3   |
| 15 | Can Deposition at the beginning of use (mg)        | 0.5   | 0.4   | 0.3   |
| 20 | Can Deposition at the end of use (mg)              | 0.6   | 0.4   | 0.3   |
| 25 | Valve Deposition at the beginning of use (mg)      | 0.2   | 0.2   | 0.2   |
| 30 | Valve Deposition at the end of use (mg)            | 0.5   | 0.3   | 0.2   |
|    | Total Drug Deposition at the beginning of use (mg) | 0.7   | 0.6   | 0.5   |
|    | Total Drug Deposition at the end of use (mg)       | 1.1   | 0.7   | 0.5   |

35 Fluticasone propionate of the present invention also shows no significant increase in the drug deposition on the can and valve through the life of the inhaler. As a result, the delivery dose is consistent through the life of the inhaler as shown in the following Example.

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Example 11  
Dose Delivered Through Use

5 The dose delivered through use of the inhalers has been measured on Inhaler Type B. Doses are collected as pairs of actuations at the beginning of use (actuations 1 & 2), the middle of use (actuations 60 & 61) and the end of use (actuations 119 & 120). The doses are collected as follows: The two actuations are fired into 10 a 500 ml separating funnel (plugged at one end with cotton wool) which has a 20 litre per minute airflow pulled through it. The separating funnel is washed with acetonitrile into a 100 ml volumetric flask containing 15 50 ml of water. The resultant solution is made up to volume and assayed by HPLC.

20 Table 14 presents the Dose Delivered Through Use data for Inhaler Type B, for which the target dose to be delivered is 44 micrograms per actuation.

Table 14

Dose Delivered Through Use for Inhaler Type B

| 25 | Drug  | Beginning of use<br>(Actuations 1 & 2) | Middle of use<br>(Actuations 60 & 61) | End of use<br>(Actuations 119 & 120) |
|----|---|--|---------------------------------------|--------------------------------------|
| 30 | Conventionally crystallised propionate (micronised)         | Mean (mcg)<br>RSD (%)                  | 31.3<br>4.2                           | 35.9<br>6.5<br>9.2                   |
|    | Fluticasone propionate of the present invention (sample 22) | Mean (mcg)<br>RSD (%)                  | 44.3<br>4.2                           | 47.9<br>6.8<br>6.3                   |

35

Fluticasone propionate of the present invention shows a

dosing profile through the life of the inhaler which is consistently close to the target dose of 44 micrograms. This profile is significantly better than that of conventionally crystallised fluticasone propionate (micronised) which shows a significant increase in dose per actuation through use of the inhaler.

10 The dose variability at each point through the use of the inhaler for fluticasone propionate of the present invention is comparable with that of conventionally crystallised fluticasone propionate (micronised) but shows an improvement towards the end of the use of the inhaler.

15 The delivered dose for fluticasone propionate of the present invention is consistently higher than that of conventionally crystallised fluticasone propionate (micronised), due to the lower drug deposition on the can and valve as shown in Example 10.

20

Example 12

Cascade Impactor Test

25 Cascade Impaction testing was performed on Inhaler Type A. The method used was in accordance with "Preparations for Inhalation; Aerodynamic assessment of fine particles using apparatus D" as defined in the British Pharmacopoeia 1993, Addendum, 1996, page A527 as applied to a metered dose inhaler formulation.

30

Data for Cascade impaction are presented in Table 15 below for a batch of inhalers made using the material from sample 19 and compared to inhalers made using conventionally crystallised and micronised material.

35

The deposition in stages 3, 4 and 5 represents the fine particle mass reaching the deep lung. The delivered

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dose (ex actuator) represents the total dose available for inhalation and the emptying efficiency of the drug from the device.

Table 15

5

## Cascade Impaction Data for Inhaler Type A

|    |                                  | Quantity of Drug Deposited on Each Stage of the Cascade Impactor (micrograms) |   |
|----|----------------------------------|---|---|
|    |                                  | Conventionally crystallised fluticasone propionate (micronised)               | Fluticasone propionate of the present invention (sample 19) |
| 10 | Mean Particle Size (microns)     | 1.8   | 4.8   |
|    | Actuator                         | 13.6  | 13.8  |
|    | Throat                           | 46.8  | 42.2  |
|    | Stage 0                          | 5.7   | 10.9  |
|    | Stage 1                          | 2.4   | 3.7   |
|    | Stage 2                          | 3.8   | 3.9   |
|    | Stage 3                          | 15.3  | 11.5  |
|    | Stage 4                          | 19.2  | 17.1  |
|    | Stage 5                          | 11.5  | 18.0  |
|    | Stage 6                          | 1.2   | 2.7   |
| 15 | Stage 7                          | 0.4   | 0.6   |
|    | Filter                           | 0.3   | 0.3   |
|    | Delivered dose Ex Valve          | 119.9   | 124.5   |
| 20 | Delivered dose Ex Actuator       | 106.3   | 110.8   |
|    | Fine Particle Mass               | 46.0  | 46.6  |
|    | Mass Median Aerodynamic diameter | 3.1   | 3.0   |
| 25 | Geometric standard deviation     | 1.7   | 1.9   |

Fluticasone propionate of the present invention shows no significant improvement in fine particle mass. The interesting feature of the present invention is that the 5 supercritical fluid crystallised fluticasone propionate with a particle size greater than that of conventionally crystallised fluticasone propionate (micronised) gives an equivalent fine particle mass in the Cascade Impactor.

10 Fluticasone propionate of the present invention shows an improved delivered dose indicating that the drug is emptied well from the device and is presenting a greater quantity of drug for inhalation. This is being aided by 15 the lower drug can/valve deposition, which increases the concentration of drug in the suspension. Again, the interesting feature of the present invention is that the supercritical fluid crystallised fluticasone propionate with a particle size greater than that of conventionally 20 crystallised fluticasone propionate (micronised) gives a greater delivered dose.

25 The data indicate that fluticasone propionate of the present invention has improved fluidisability and flow properties.

CLAIMS:

1. S-fluoromethyl 6 $\alpha$ ,9 $\alpha$ -difluoro-11 $\beta$ -hydroxy-16 $\alpha$ -methyl-3-oxo-17 $\alpha$ -propionyloxyandrosta-1,4-diene-17 $\beta$ -carbothiate (fluticasone propionate) in an easily handled and easily fluidised crystalline form, with a controlled particle size and shape.
2. S-fluoromethyl 6 $\alpha$ ,9 $\alpha$ -difluoro-11 $\beta$ -hydroxy-16 $\alpha$ -methyl-3-oxo-17 $\alpha$ -propionyloxyandrosta-1,4-diene-17 $\beta$ -carbothiate (fluticasone propionate) having an orthorhombic crystalline structure.
3. Fluticasone propionate as claimed in claim 1 or claim 2 in a form with a dynamic bulk density of less than 0.2 g cm<sup>-3</sup>.
4. Fluticasone propionate as claimed in claim 3 with a dynamic bulk density in the range between 0.05 and 0.17 g cm<sup>-3</sup>.
5. Fluticasone propionate as claimed in claim 4 with a dynamic bulk density in the range between 0.05 and 0.08 g cm<sup>-3</sup>.
6. Fluticasone propionate as claimed in any one of claims 1 to 5 which has a particle size in the range of 1 to 10 microns.
7. Fluticasone propionate as claimed in any one of claims 1 to 6 which has a uniformity coefficient of from 1 to 20.
8. Fluticasone propionate as claimed in any one of claims 1 to 7 which has a respirable fraction of 14% or more by weight.

- 50 -

9. Fluticasone propionate as claimed in any one of claims 1 to 8 which has a cohesivity of 0 to 20%.

10. Fluticasone propionate as claimed in claim 9 which 5 has a cohesivity of 0 to 10%.

11. Fluticasone propionate as claimed in any one of claims 1 to 10 being pure polymorphic Form 2 characterised by an X-ray powder diffraction trace 10 substantially as shown in Figure 23.

12. A process for preparing fluticasone propionate as claimed in any one of claims 1 to 11, comprising the co-introduction of a supercritical fluid and a vehicle 15 containing at least fluticasone propionate in solution or suspension into a particle formation vessel, the temperature and pressure in which are controlled, such that dispersion and extraction of the vehicle occur substantially simultaneously by the action of the 20 supercritical fluid.

13. A process as claimed in claim 12 wherein the co-introduction of the supercritical fluid and the vehicle containing at least fluticasone propionate in solution 25 or suspension is effected using a nozzle of coaxial design.

14. A process as claimed in claim 12 or claim 13 wherein the supercritical fluid is carbon dioxide.

30 15. A pharmaceutical composition comprising fluticasone propionate as claimed in any one of claims 1 to 11 together with at least one pharmaceutically acceptable carrier or excipient.

35 16. A pharmaceutical composition as claimed in claim 15 wherein the carrier is silicon dioxide or hydroxypropyl-

cellulose.

17. A pharmaceutical composition as claimed in claim 15 wherein the carrier is lactose.

5

18. A pharmaceutical composition as claimed in claim 17 in the form of a dry powder suitable for inhalation.

10 19. A pharmaceutical composition as claimed in any one of claims 15 to 17 in the form of an aerosol spray presentation.

15 20. A pharmaceutical composition as claimed in claim 19 in which the aerosol spray presentation is a metered dose inhaler.

20 21. A pharmaceutical composition as claimed in claim 19 or claim 20 wherein the aerosol spray presentation comprises HFA134a as propellant.

25

22. A pharmaceutical composition as claimed in any one of claims 15 to 21 comprising multicomponent particles comprising fluticasone propionate and carrier.

30

23. A process for preparing a pharmaceutical composition as claimed in claim 22, comprising the co-introduction of a supercritical fluid, the carrier and a vehicle containing at least fluticasone propionate in solution or suspension into a particle formation vessel, the temperature and pressure in which are controlled, such that dispersion and extraction of the vehicle occur substantially simultaneously by the action of the supercritical fluid, and the fluticasone propionate and carrier are co-crystallised together.

35

24. A process as claimed in claim 23 wherein the co-introduction of the supercritical fluid, the carrier and

the vehicle containing at least fluticasone propionate in solution or suspension is effected using a nozzle of coaxial design.

5 25. Fluticasone propionate as claimed in any one of claims 1 to 11 for use in therapy.

10 26. Use of fluticasone propionate as claimed in any one of claims 1 to 11 for the manufacture of a medicament for combating respiratory disorders.

27. Use as claimed in claim 26 wherein the respiratory disorder is asthma.

15 28. A method of treatment of a human or animal body suffering from a respiratory disorder, comprising administering to said human or animal body an effective amount of fluticasone propionate as claimed in any one of claims 1 to 11.

20 29. A method of treatment as claimed in claim 28 wherein the respiratory disorder is asthma.

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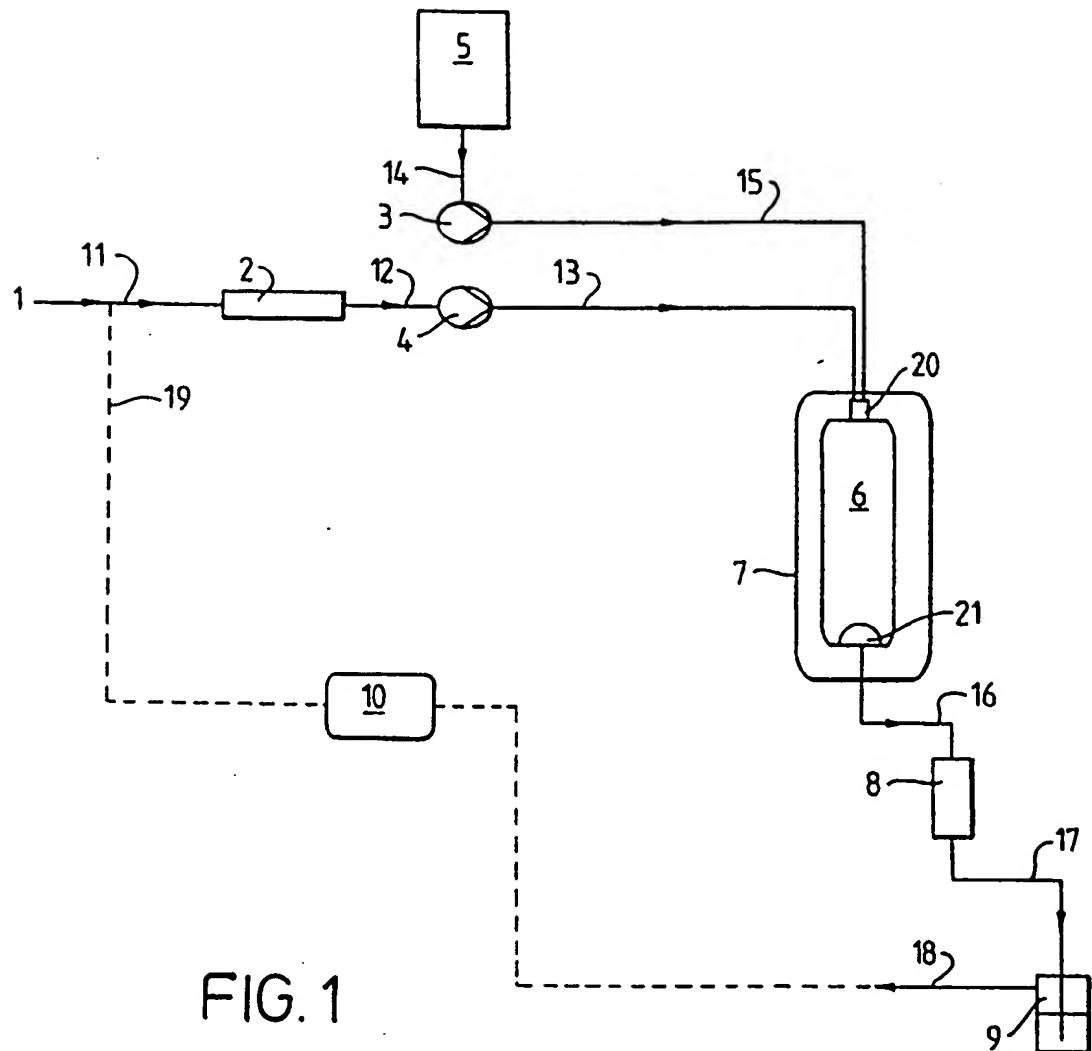


FIG. 1

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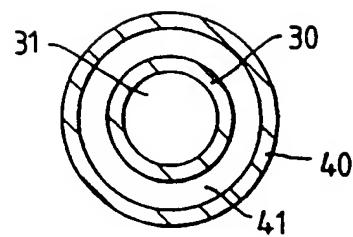


FIG. 2A

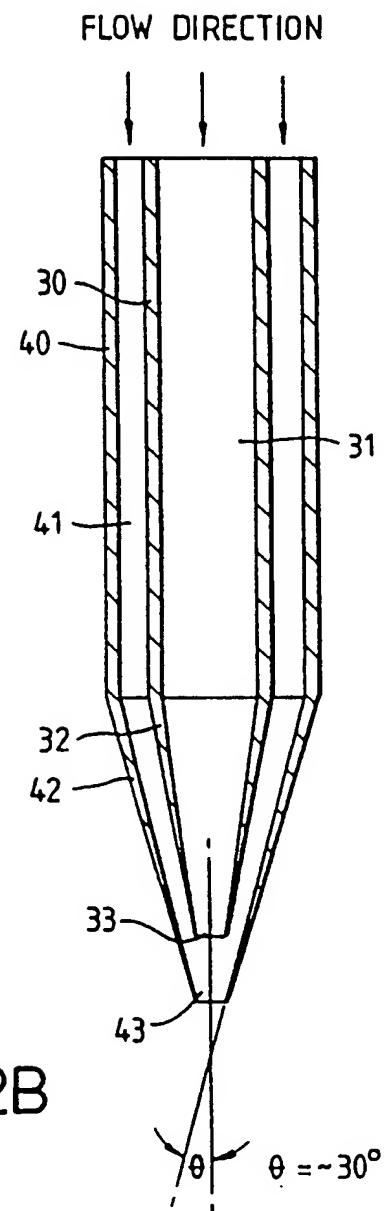
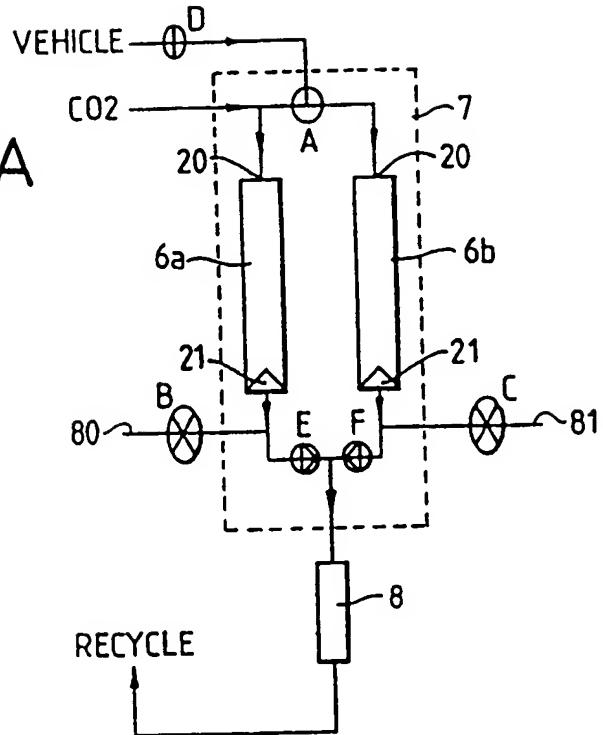


FIG. 2B

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FIG. 3A



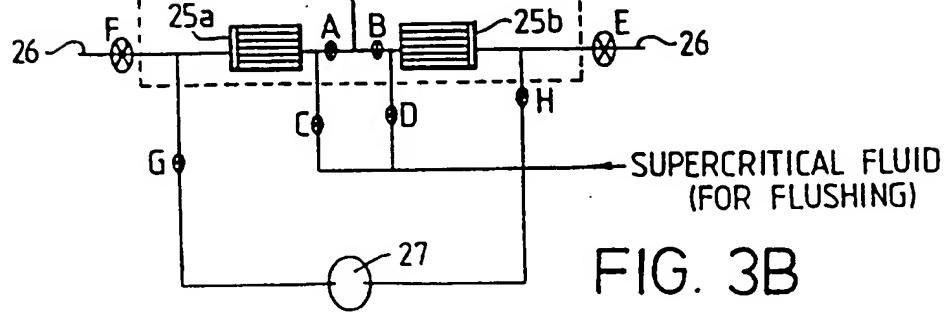
SUPERCritical FLUID

SOLUTION

20

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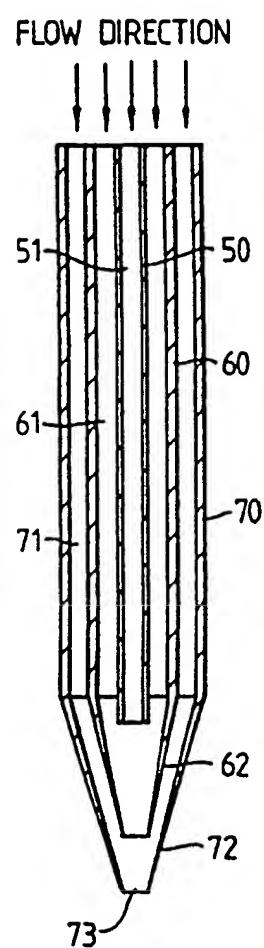


FIG. 4

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FIG. 5

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FIG. 6

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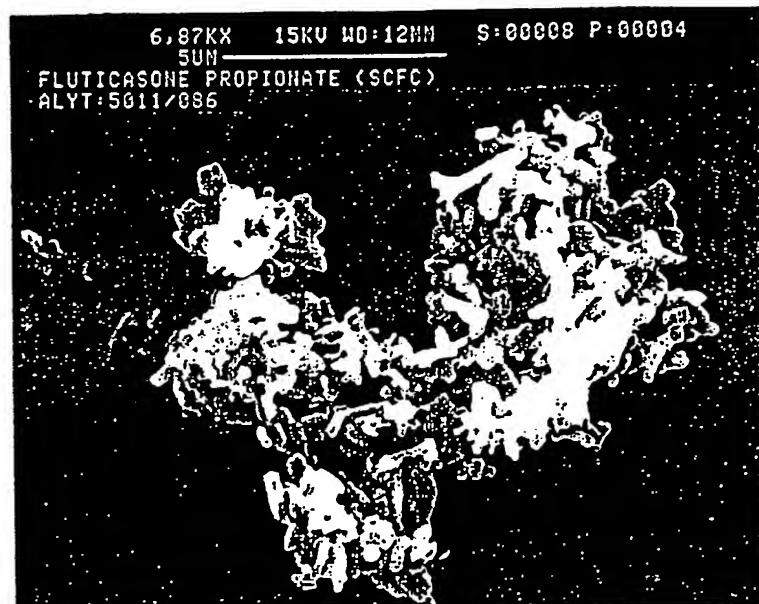
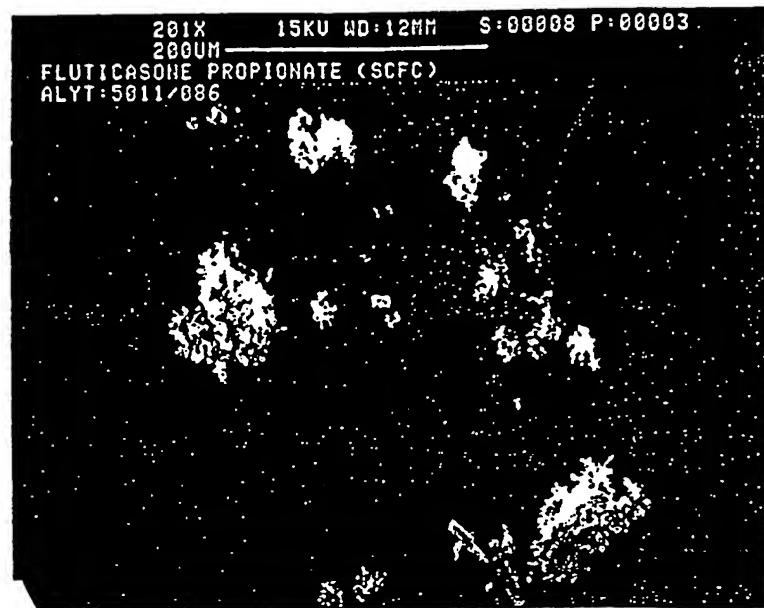


FIG. 7

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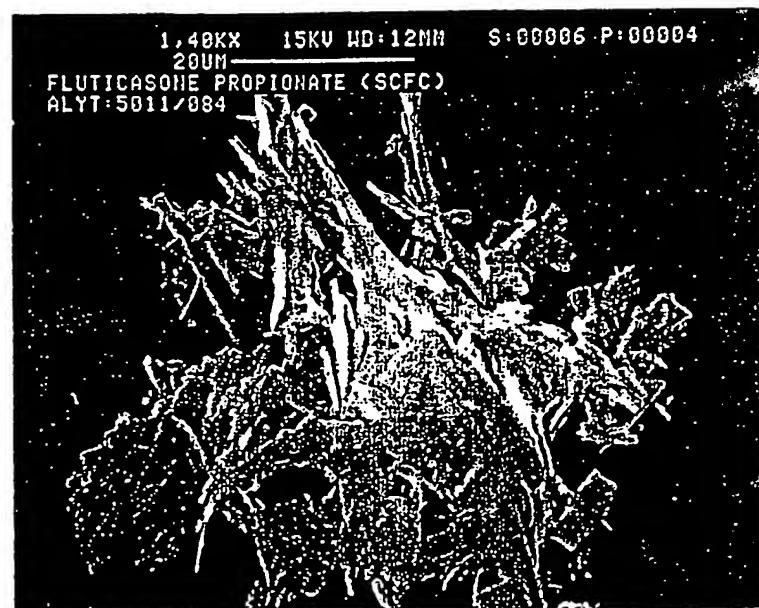
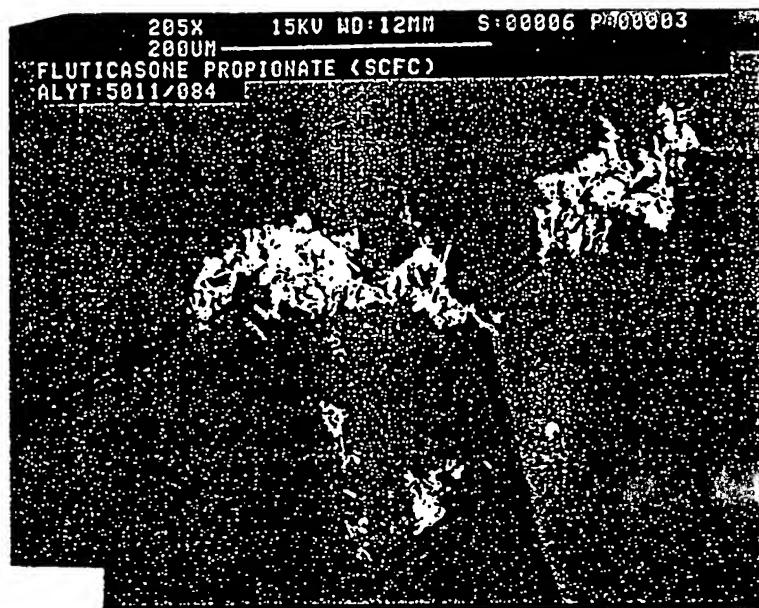
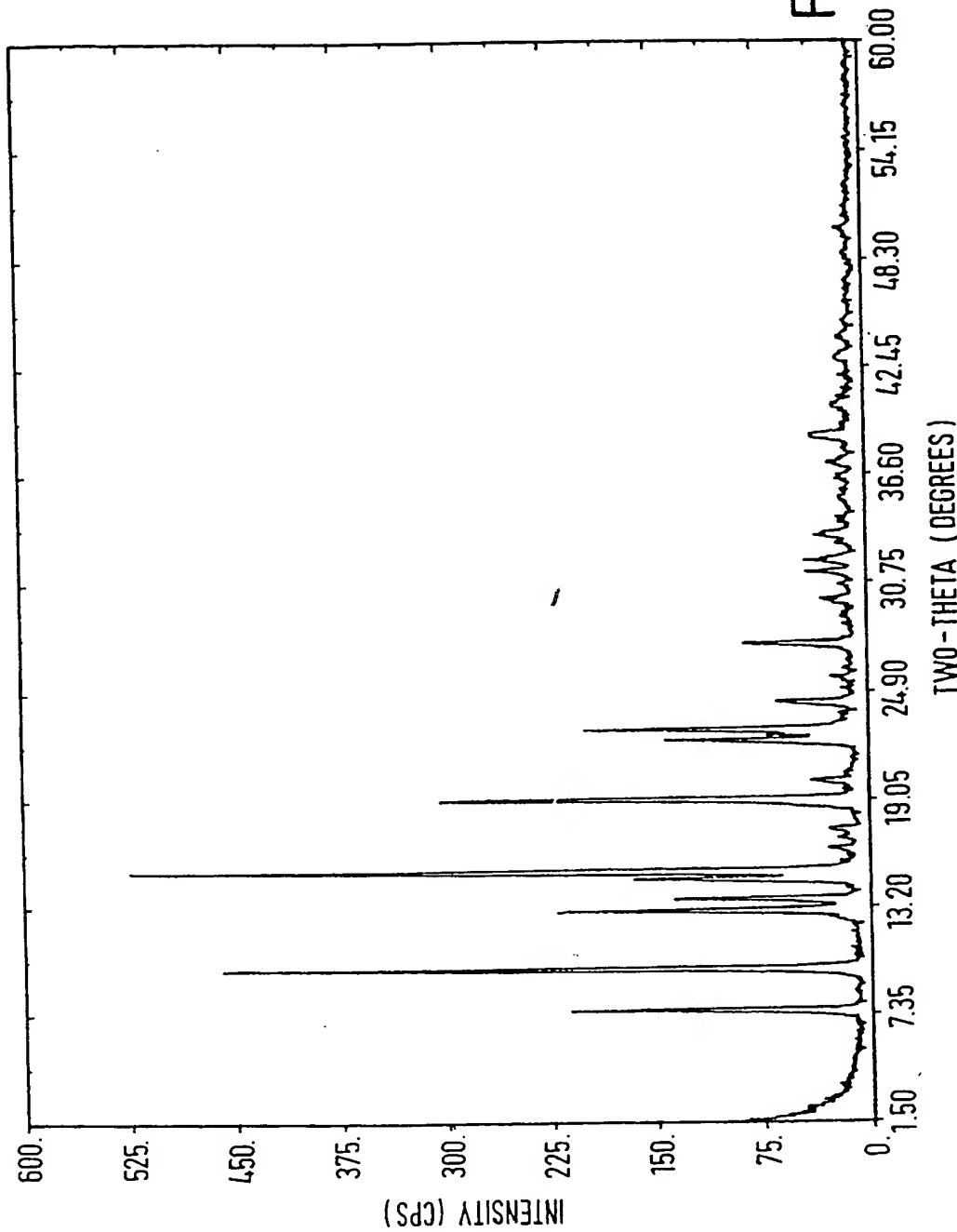


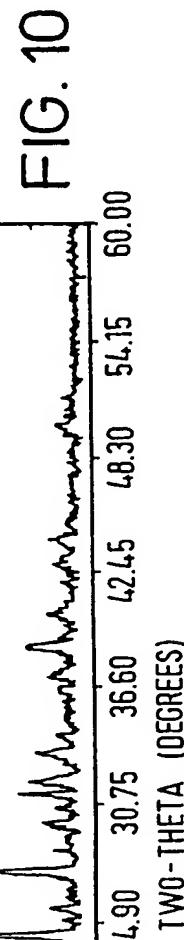
FIG. 8

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FIG. 9



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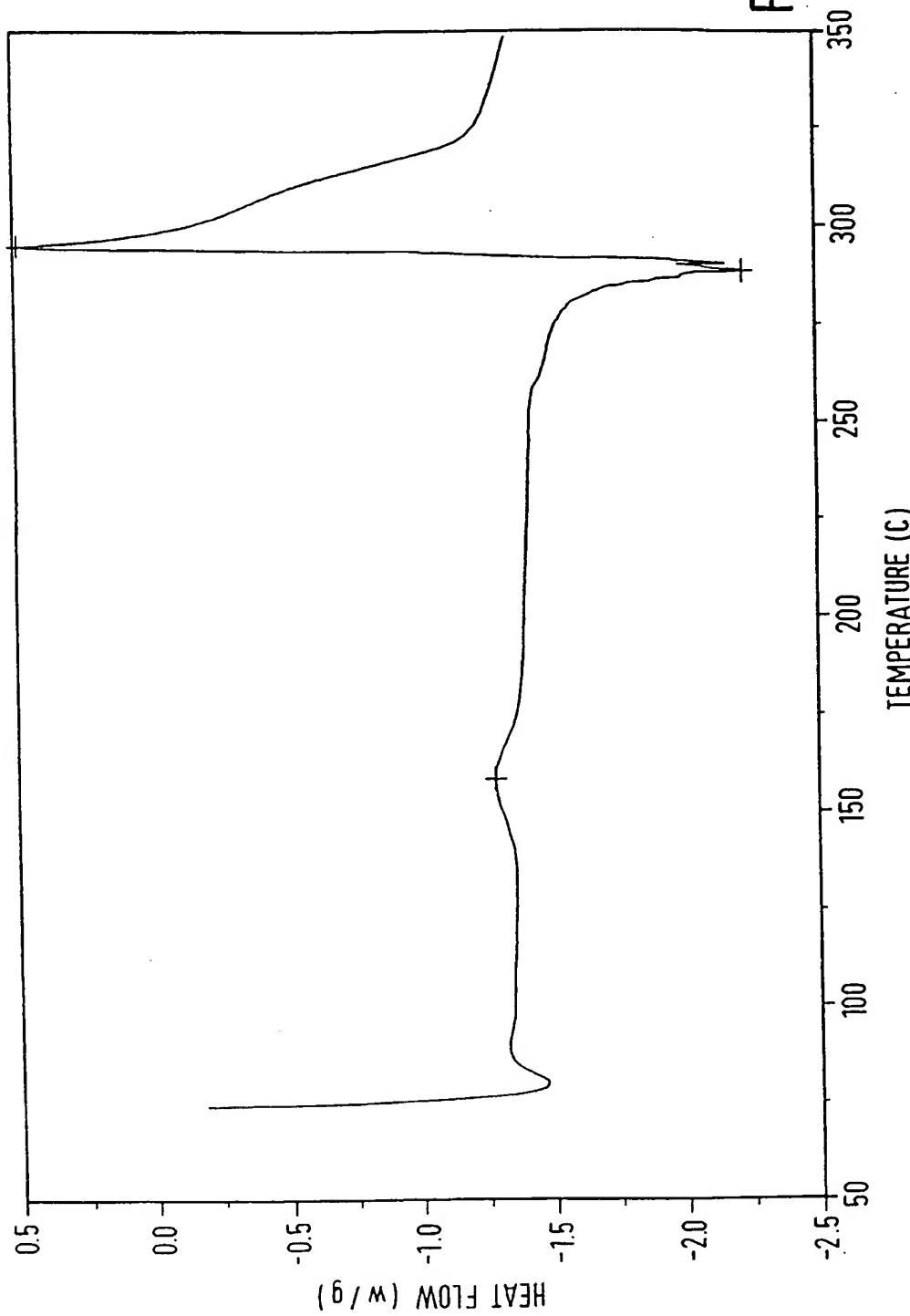


137.5  
375.0  
312.5  
250.0  
187.5  
125.0  
62.5  
0.0

INTENSITY (CPS)

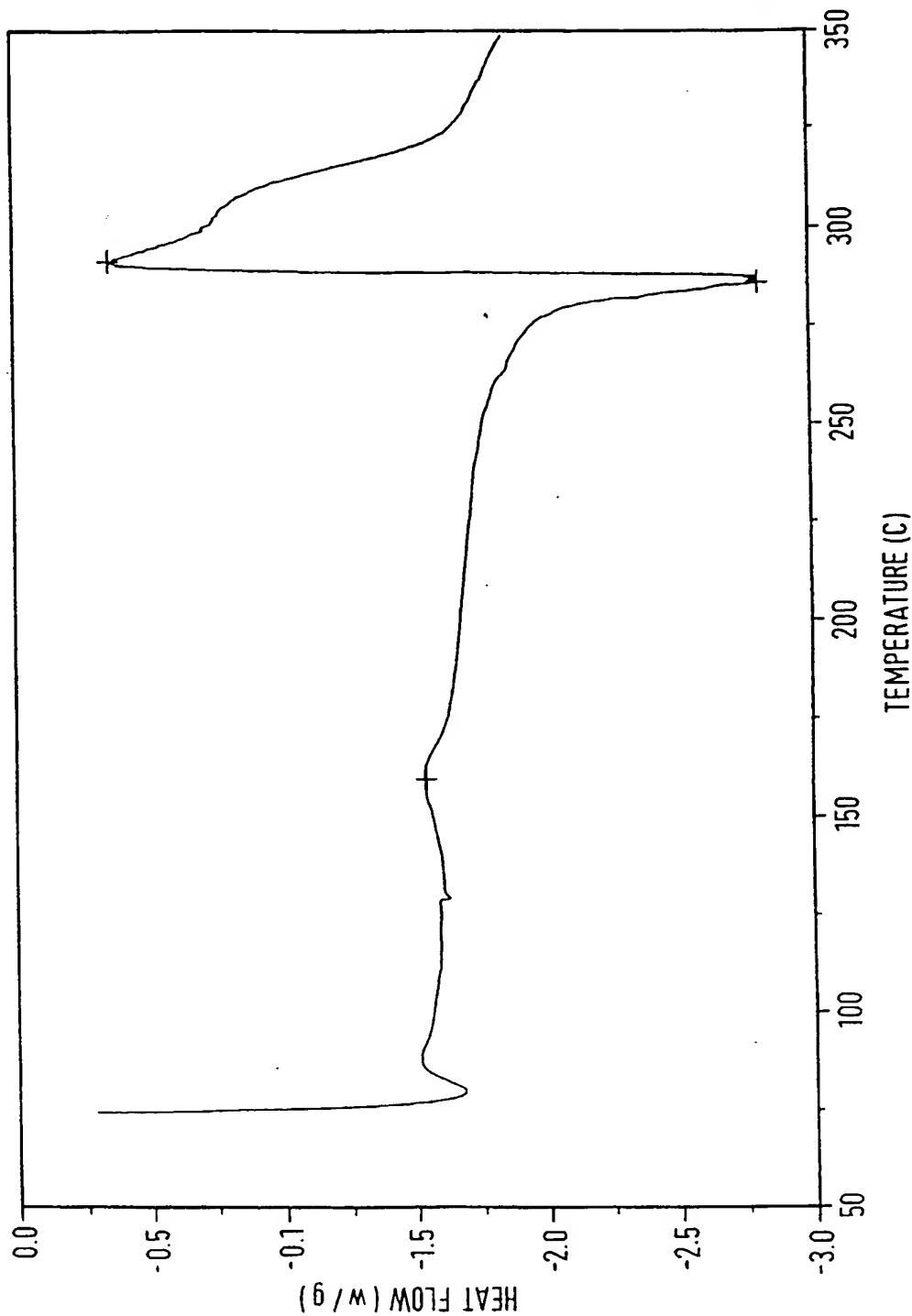
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FIG. 11



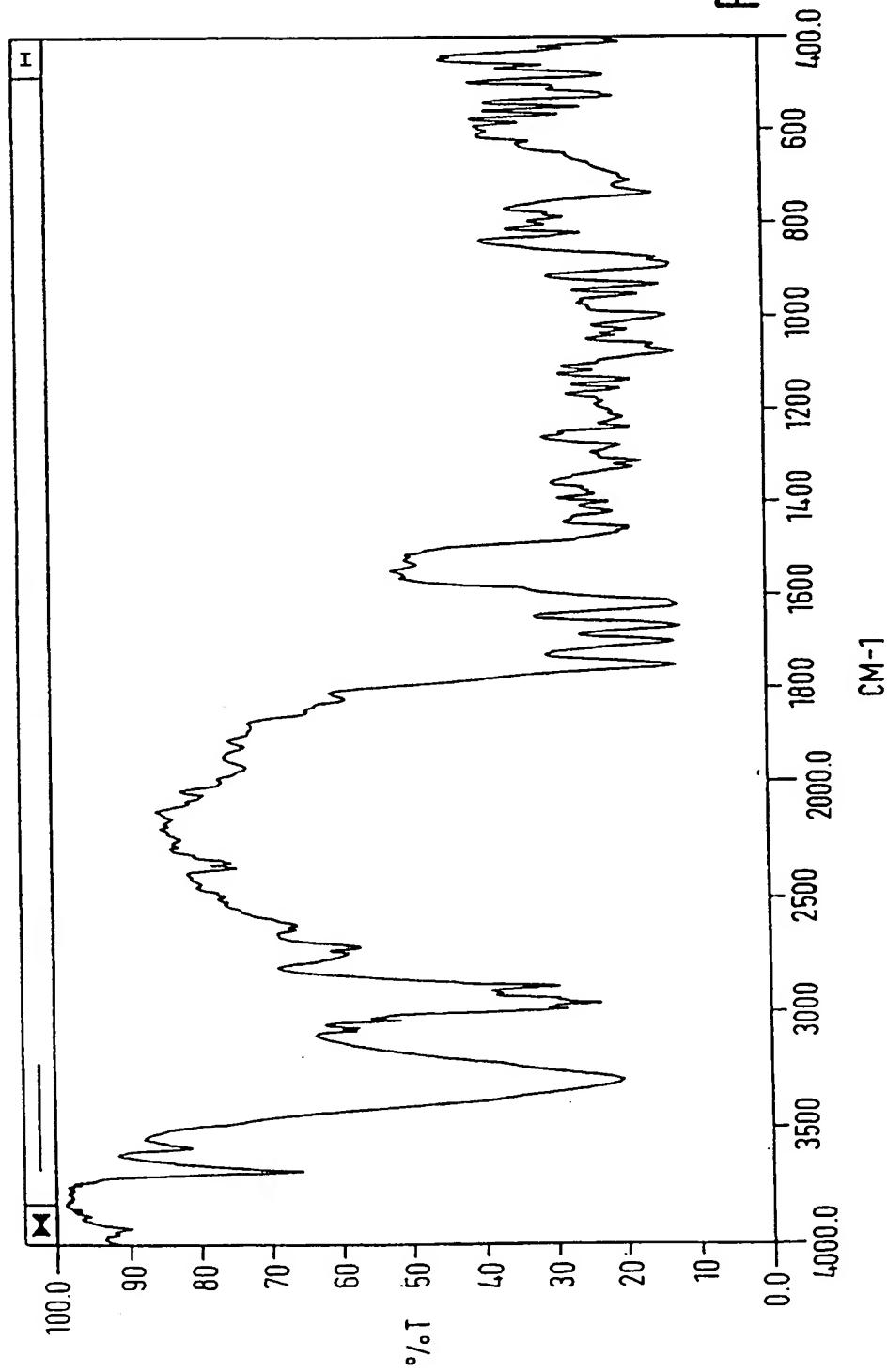
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FIG. 12



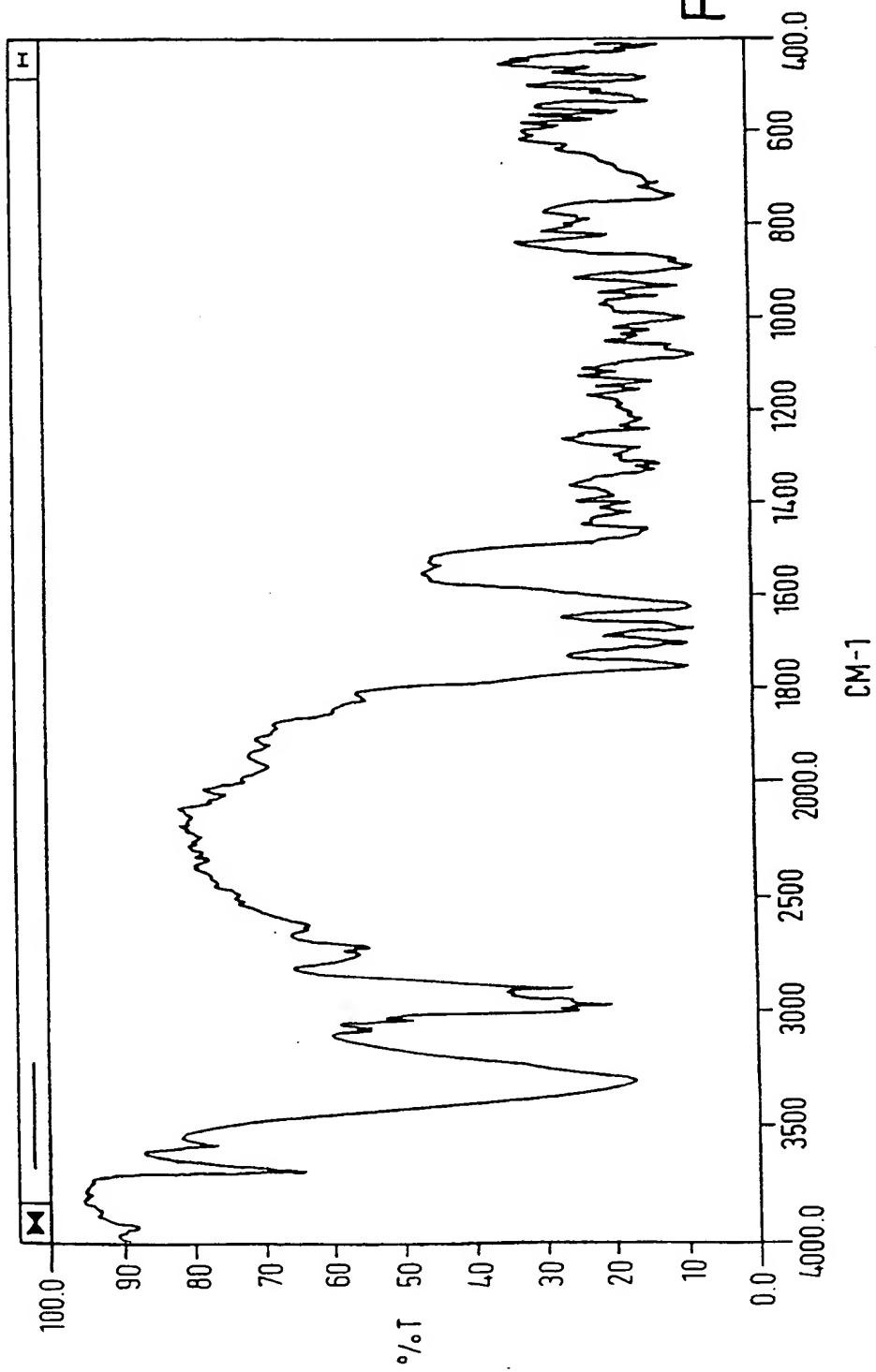
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FIG. 13



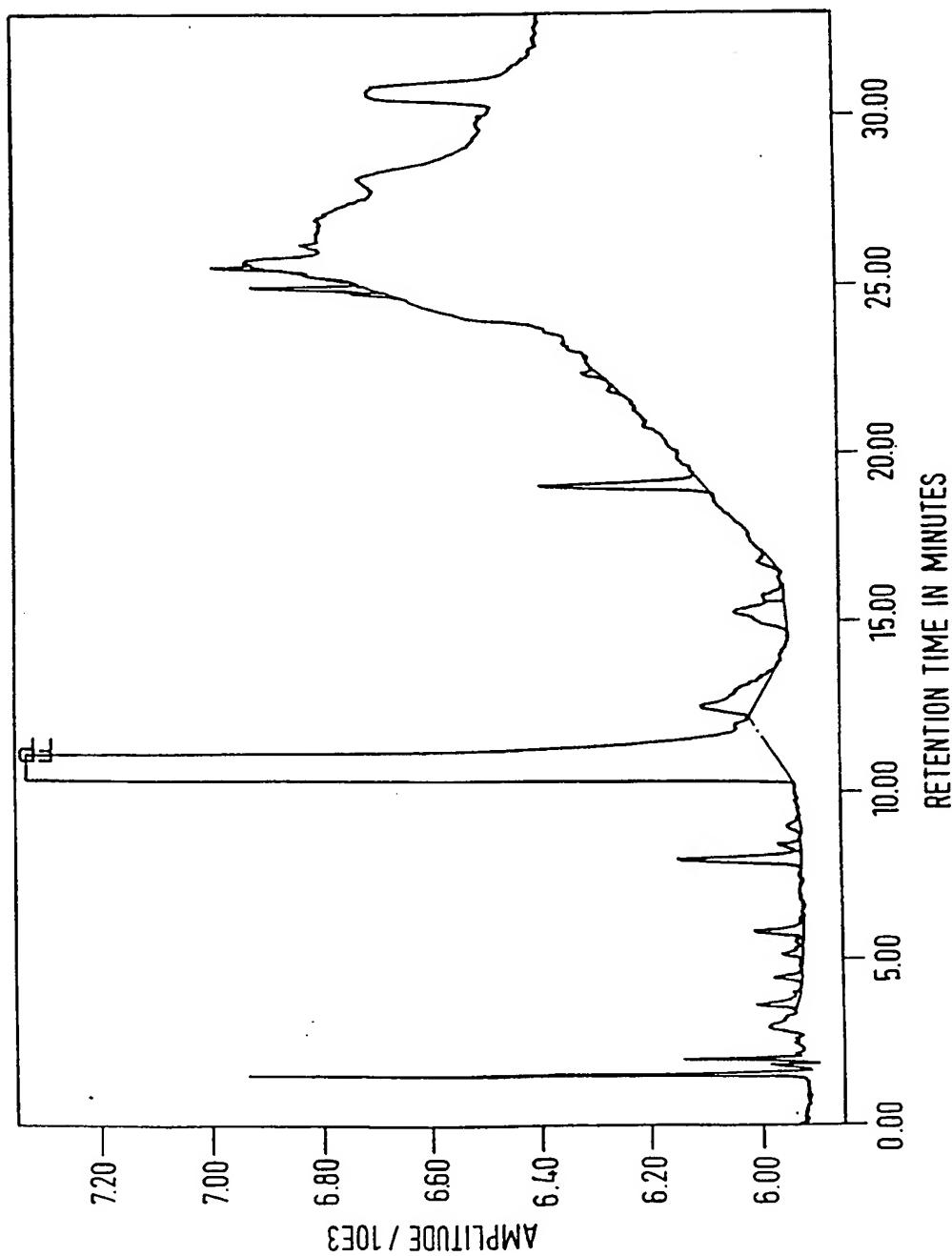
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FIG. 14



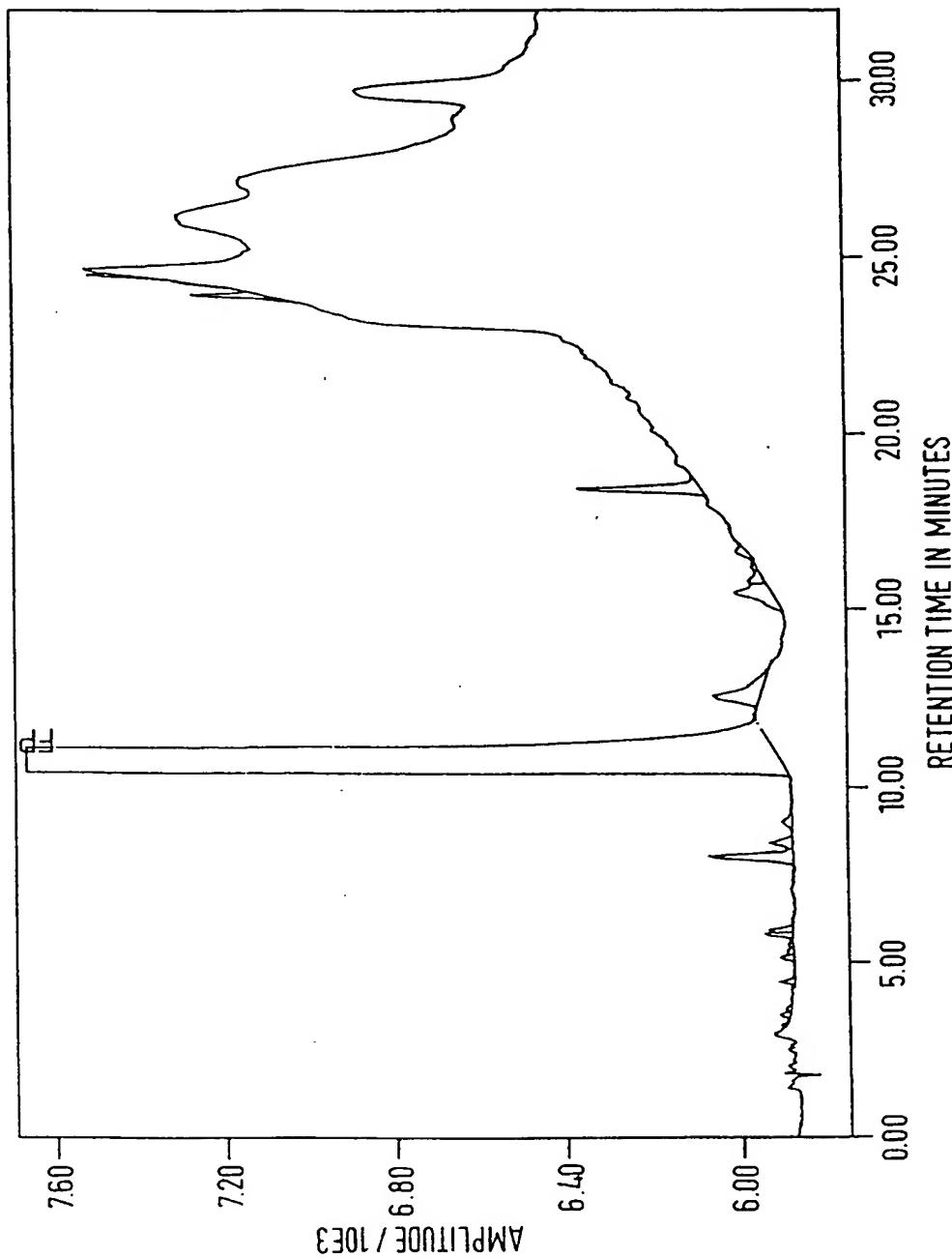
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FIG. 15



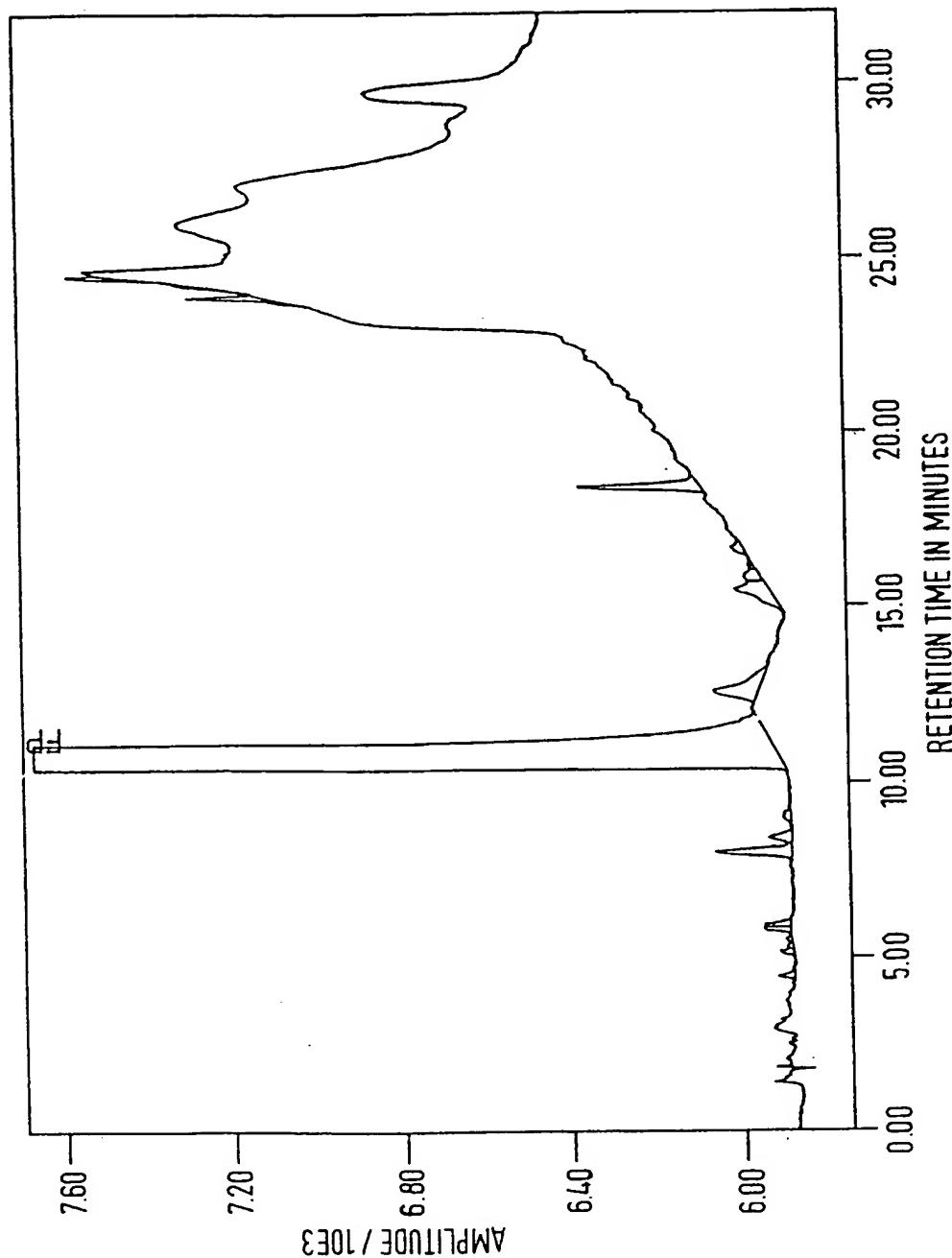
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FIG. 16



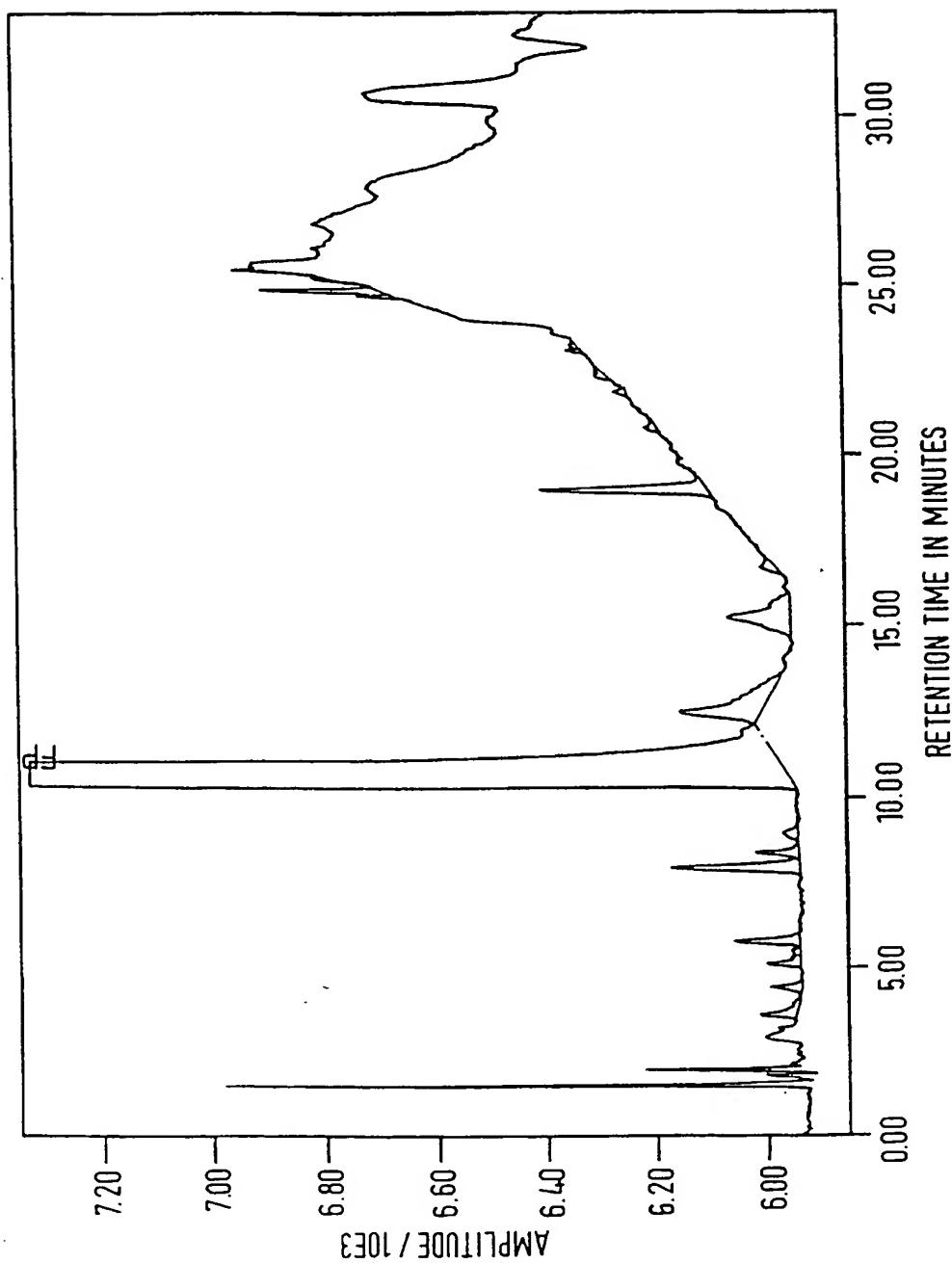
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FIG. 17



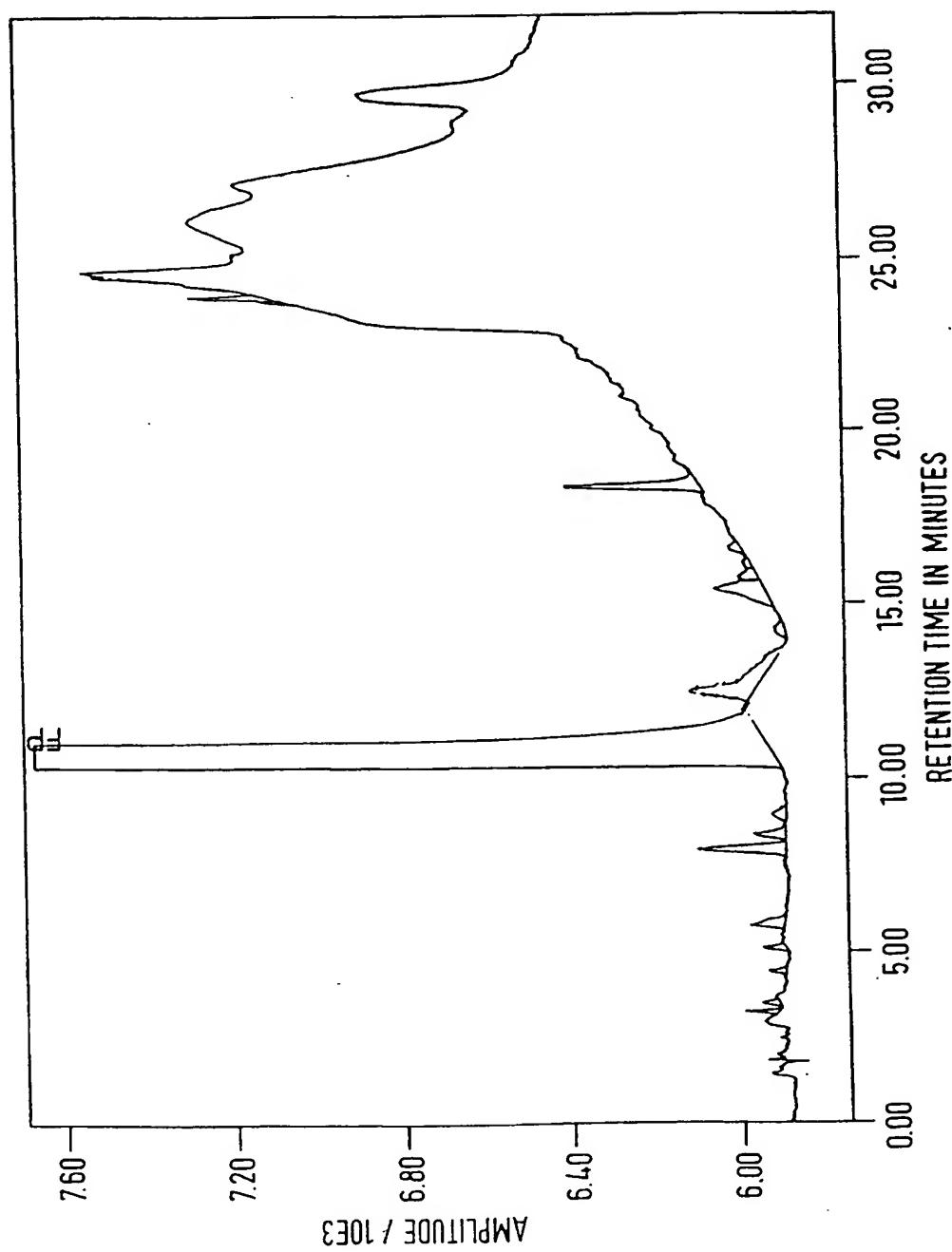
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FIG. 18



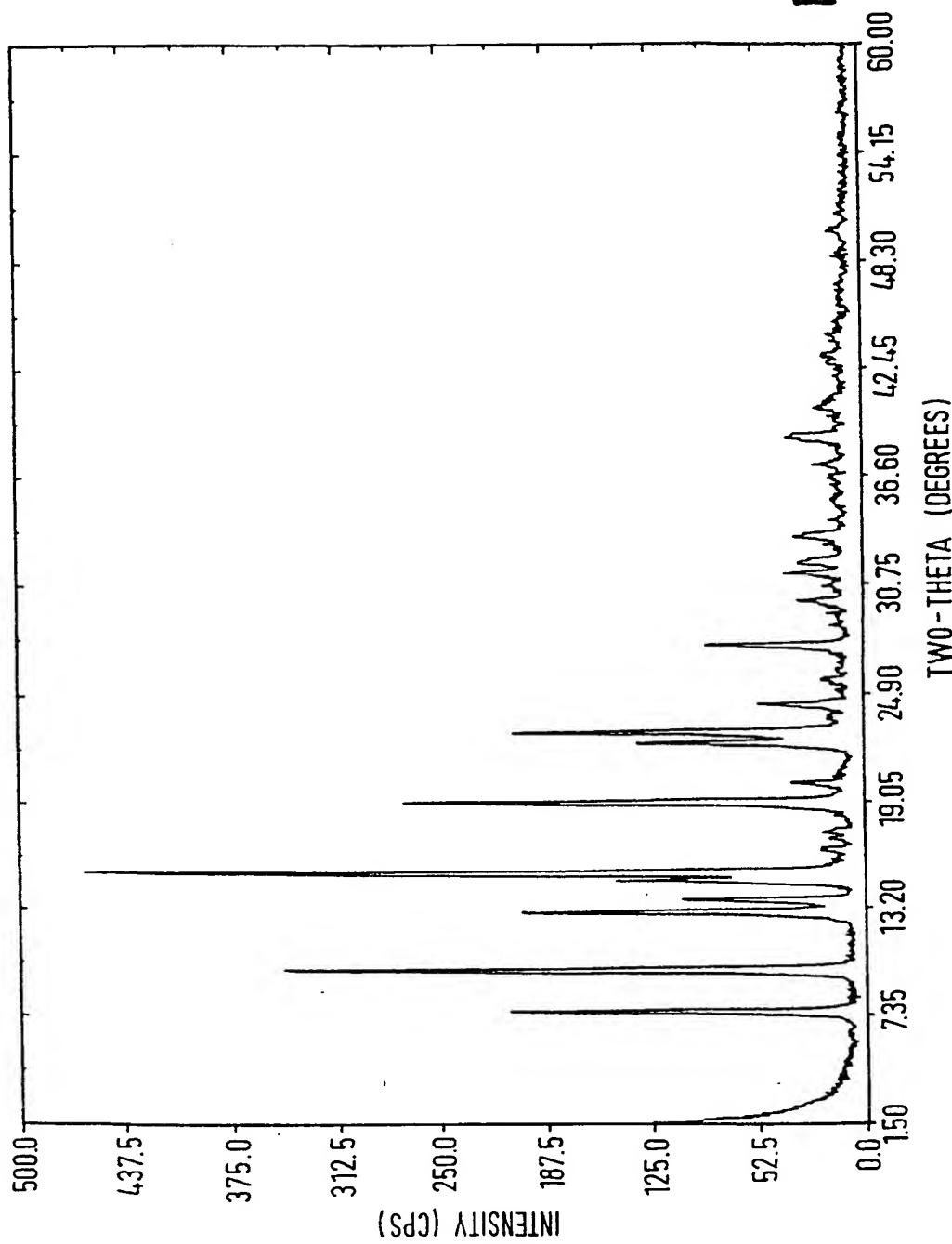
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FIG. 19



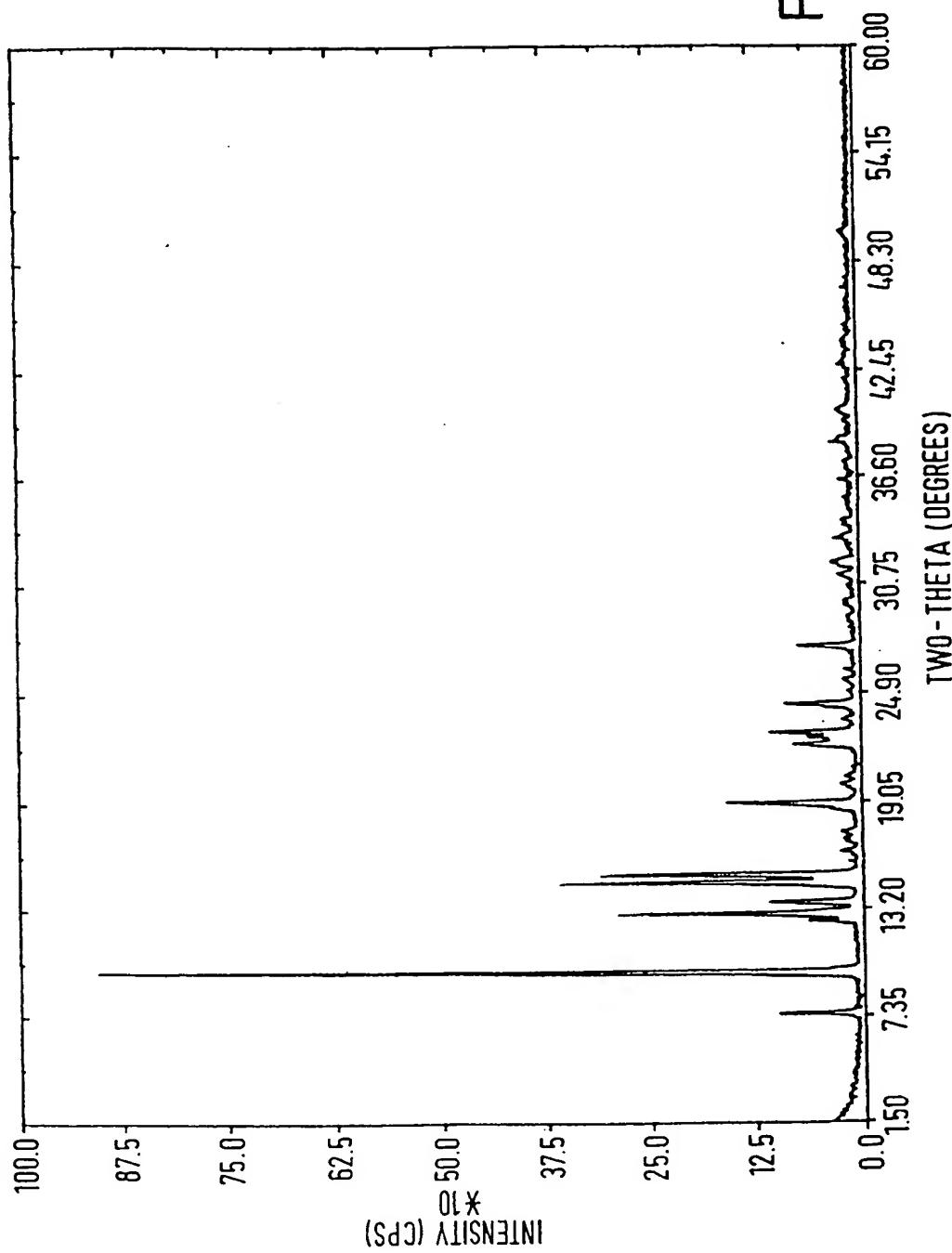
20 / 24

FIG. 20



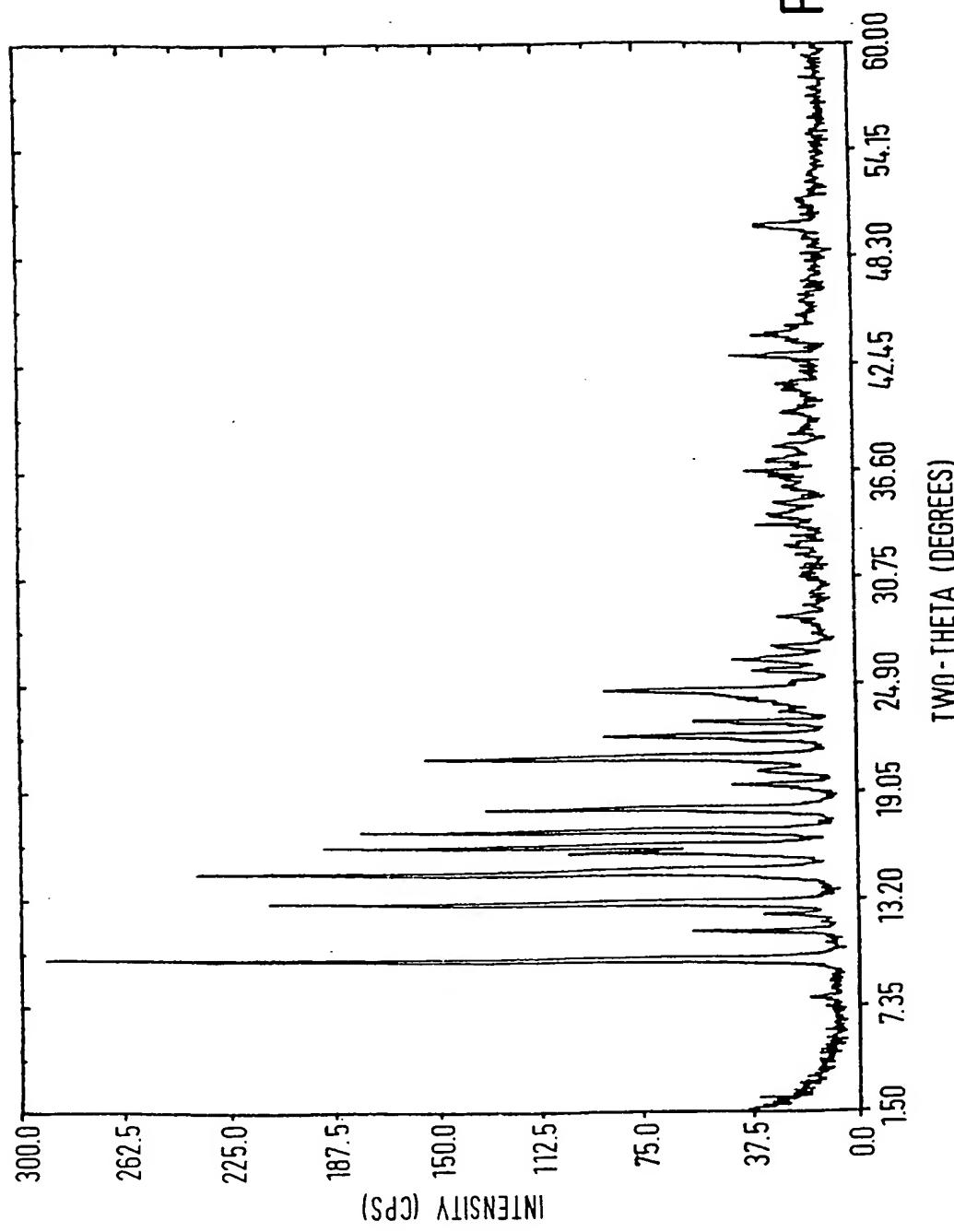
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FIG. 21



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FIG. 22



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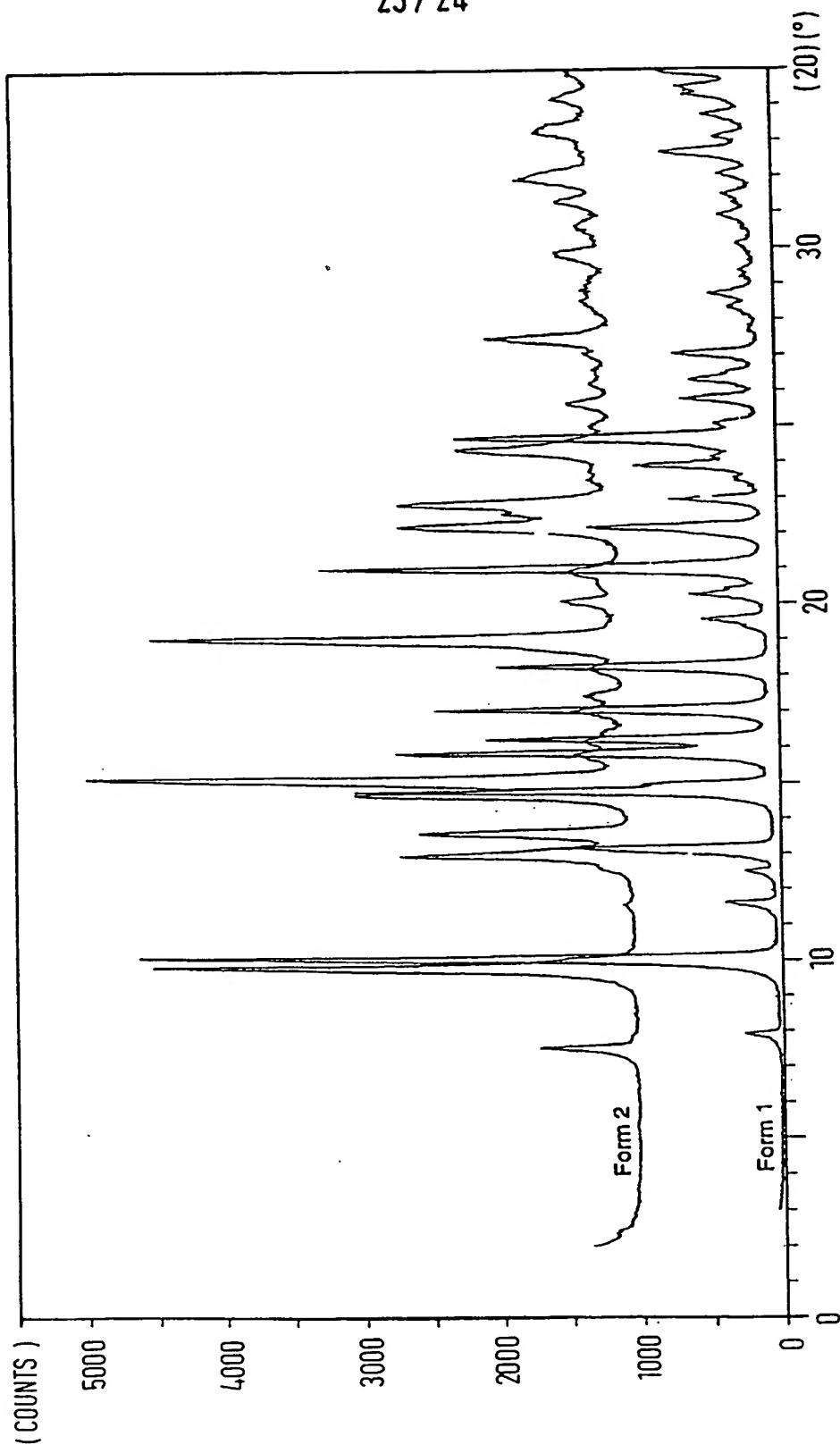


FIG. 23

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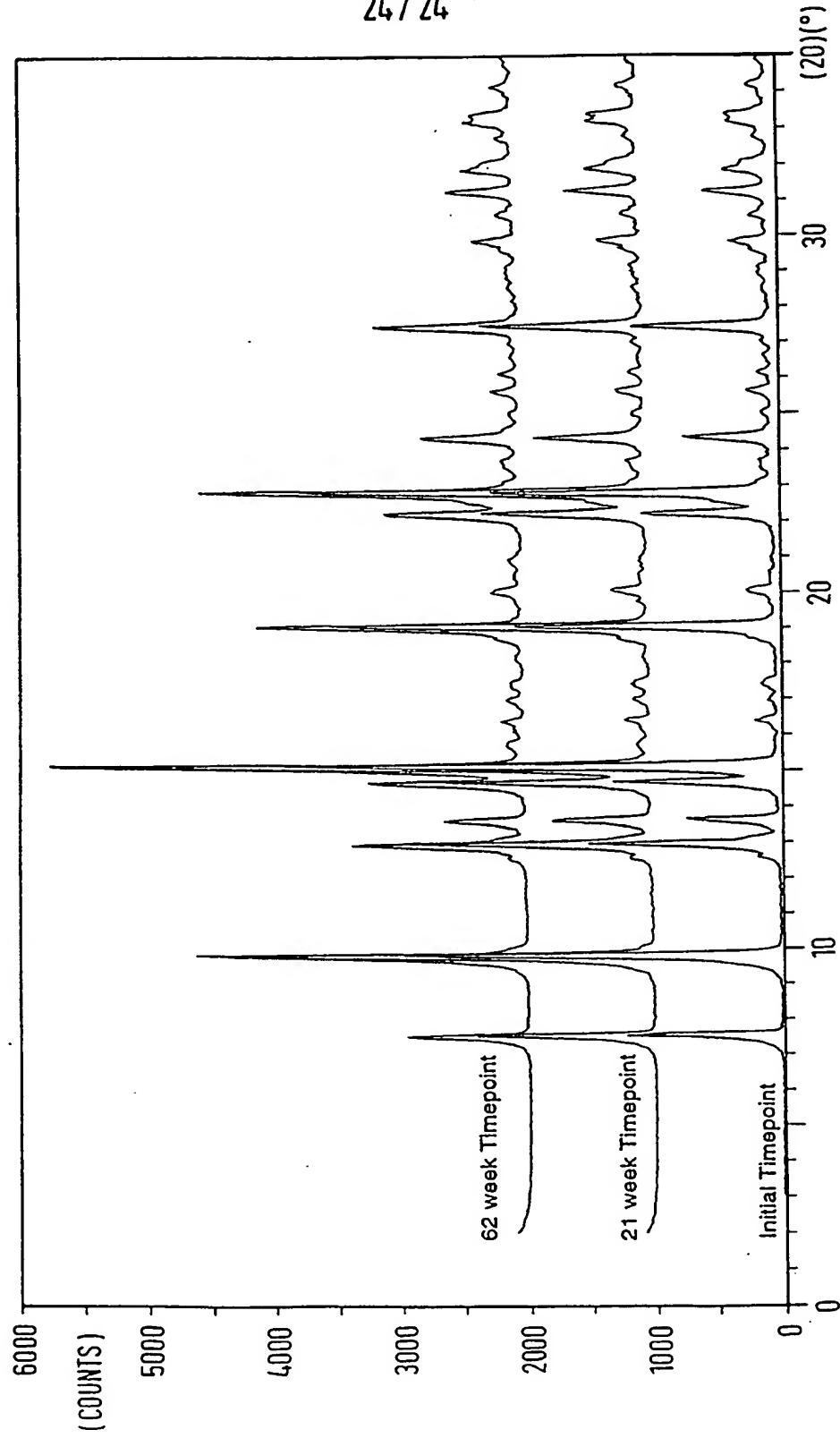


FIG. 24

# INTERNATIONAL SEARCH REPORT

Int. Application No

PCT/GB 97/02929

**A. CLASSIFICATION OF SUBJECT MATTER**  
 IPC 6 C07J31/00 A61K31/56

According to International Patent Classification(IPC) or to both national classification and IPC

**B. FIELDS SEARCHED**

Minimum documentation searched (classification system followed by classification symbols)  
 IPC 6 C07J A61K

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

**C. DOCUMENTS CONSIDERED TO BE RELEVANT**

| Category | Citation of document, with indication, where appropriate, of the relevant passages  | Relevant to claim No. |
|----------|---|-----------------------|
| A        | GB 2 088 877 A (GLAXO GROUP LTD) 16 June 1982<br>cited in the application<br>see example 19<br>---  | 2-29                  |
| A        | PHILLIPPS G H ET AL: "Synthesis and Structure-Activity Relationships in a Series of Antiinflammatory Corticosteroid Analogs, Halomethyl Androstan-17.beta.-carbothioates and -17.beta.-carboseleenoates"<br>JOURNAL OF MEDICINAL CHEMISTRY.,<br>vol. 37, no. 22, 28 October 1994,<br>WASHINGTON US,<br>pages 3717-3729, XP002052064<br>see page 3726, column 2; example 13E<br>---<br>-/- | 2-29                  |

Further documents are listed in the continuation of box C.

Patent family members are listed in annex.

Special categories of cited documents :

- "A" document defining the general state of the art which is not considered to be of particular relevance
- "E" earlier document but published on or after the international filing date
- "L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- "O" document referring to an oral disclosure, use, exhibition or other means
- "P" document published prior to the international filing date but later than the priority date claimed

"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

"X" document of particular relevance: the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

"Y" document of particular relevance: the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.

"&" document member of the same patent family

1

Date of the actual completion of the international search

14 January 1998

Date of mailing of the international search report

03.02.98

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Authorized officer

Watchorn, P

## INTERNATIONAL SEARCH REPORT

International Application No  
PCT/GB 97/02929

## C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

| Category | Citation of document, with indication, where appropriate, of the relevant passages  | Relevant to claim No. |
|----------|---|-----------------------|
| P,X      | STECKEL H ET AL: "Micronizing of steroids for pulmonary delivery by supercritical carbon dioxide"<br>INT. J. PHARM. ,<br>vol. 152, no. 1, 1997,<br>pages 99-110, XP002052065<br>see the whole document<br>----- | 2-29                  |

## INTERNATIONAL SEARCH REPORT

I national application No.

PCT/GB 97/02929

### Box I Observations where certain claims were found unsearchable (Continuation of item 1 of first sheet)

This International Search Report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1.  Claims Nos.: because they relate to subject matter not required to be searched by this Authority, namely:

Remark: Although claim(s) 28 and 29 is(are) directed to a method of treatment of the human/animal body, the search has been carried out and based on the alleged effects of the compound/composition.

2.  Claims Nos.: because they relate to parts of the International Application that do not comply with the prescribed requirements to such an extent that no meaningful International Search can be carried out, specifically:

See FURTHER INFORMATION sheet PCT/ISA/210

3.  Claims Nos.: because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

### Box II Observations where unity of invention is lacking (Continuation of item 2 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

1.  As all required additional search fees were timely paid by the applicant, this International Search Report covers all searchable claims.

2.  As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.

3.  As only some of the required additional search fees were timely paid by the applicant, this International Search Report covers only those claims for which fees were paid, specifically claims Nos.:

4.  No required additional search fees were timely paid by the applicant. Consequently, this International Search Report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

#### Remark on Protest

The additional search fees were accompanied by the applicant's protest.

No protest accompanied the payment of additional search fees.

FURTHER INFORMATION CONTINUED FROM PCT/ISA/ 210

Claims Nos.: 1 (in full), 3-11,15-29 (in part)

because they relate to parts of the international application that do not comply with the prescribed requirements to such an extent that no meaningful international search can be carried out, specifically:

Claim 1 defines the crystalline products of the present application in terms of a result to be achieved, i.e. in terms of the solution to the problem of providing easily handled, easily fluidised crystals of Fluticasone propionate (FP) with controlled particle size and shape, without giving the skilled person any indication whatsoever as to the technical means to achieve solve this problem. Claim 1 is consequently so unclear according to Article 6 PCT, that no search is possible for this claim (see also the preliminary PCT examination Guidelines III 4.7). Consequently claims 3-11 which depend in part on claim 1 are not searched in as far as they depend on this claim, likewise, claims 15-29 have not been searched in as far as they relate to pharmaceutical compositions or medical uses of the products of claim 1 and the processes for the production of those compositions.

# INTERNATIONAL SEARCH REPORT

Information on patent family members

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